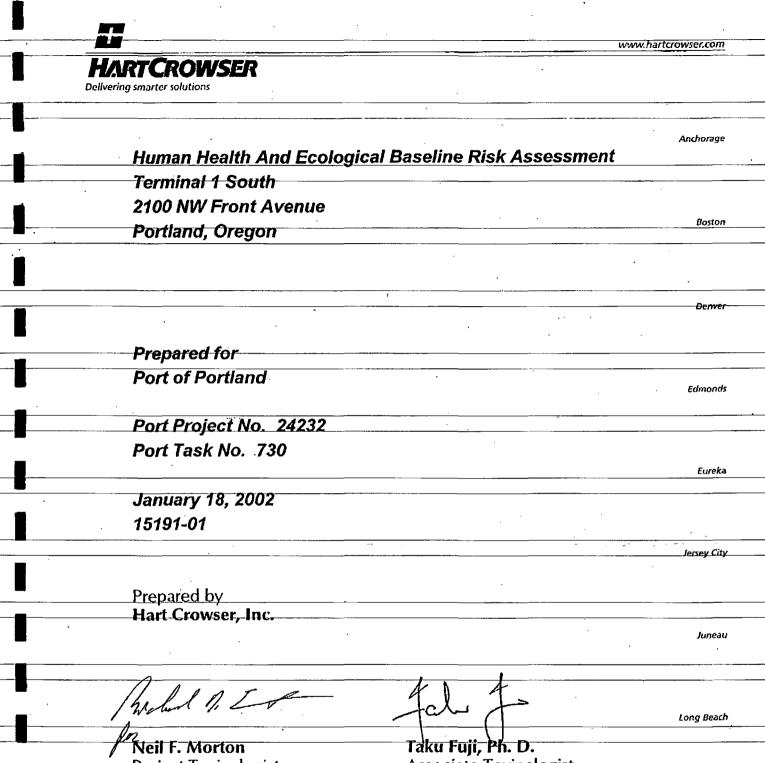
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Seattle

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ACRONY	AS	·	•
DIAL	5 LW:1.	· · · · · · · · · · · · · · · · · · ·	<u>·</u> _
BW	Body Weight		
CDI	Chronic Daily Intake		
_COPC	Chemicals of Potential Concern		
CSM	Conceptual Site Model	•	
СТ	Central Tendency		
DEQ	Oregon Department of Environmental Quality	·	
EDI	Estimated Daily Intake		
EPA	United States Environmental Protection Agency		
-EPC	Exposure Point-Concentration		
ERA	Ecological Risk Assessment		
HEAST	Health Effects Assessment Summary Table		
HHRA	Human Health Risk Assessment		
-HI	— Hazard Index	·	
HQ	Hazard Quotient		
IRIS	Integrated Risk Information System		
MS/MSD	Matrix Spike/Matrix Spike Duplicate		
NOAEL	No Observed Adverse Effect Level		
OAR	Oregon Administrative Rules		
ONHP	Oregon Natural Heritage Program		
PAH	Polycyclic Aromatic Hydrocarbon		
PORT	Port of Portland		
PRG	Preliminary Remediation Goal		
RCRA	Resource-Conservation and Recovery Act		
RfD	Reference Dose		
RI	Remedial Investigation		•
RME	Reasonable Maximum Exposure		
SF	Slope Factor		
SQL	Sample Quantitation Limit		
T&E	Threatened and Endangered	· .	_
T1S Site	Terminal 1 South Site		- · · · · · · · · · · · · · · · · · · ·
TPH	Total Petroleum Hydrocarbon	•	
UCL	Upper Confidence-Limit	•	
	• •		
-UF	Uncertainty Factor		
VOC	Volatile Organic Compound	•	
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		i,	
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HUMAN HEALTH AND ECOLOGICAL BASELINE RISK ASSESSMENT TERMINAL 1 SOUTH 2100 NW FRONT AVENUE PORTLAND, OREGON **EXECUTIVE SUMMARY** Objectives. Hart Crowser conducted a human health risk assessment (HHRA) and a Level 1 Scoping and a Modified Level 2 Screening ecological risk assessment (ERA) for the Port of Portland Terminal 1 South (T1S Site) in Portland, Oregon. The purpose of the HHRA is to evaluate potential risks and hazards to human health-associated-with each potential exposure-pathway-(complete pathways identified for the site are exposure to surface and subsurface soil and inhalation of volatile compounds from groundwater). The purpose of the Level 1 Scoping ERA is to provide a conservative, qualitative determination of whether ecological receptors and/or exposure pathways are potentially present at or in the locality of the site. The Modified Level 2 Screening ERA was conducted on site groundwater data to determine whether constituents were present at levels of concern for aquatic ecological receptors. Site Description and History. The T1S Site is located at 2100 NW Front Avenue in Portland, Oregon (Figure 1). The site consists of approximately-21 acres that are almost completely paved with asphalt or concrete or covered by buildings (Figure 2). Two primary structures, designated as Warehouse No. 2 and House No. 104, are currently located at the T15 Site. An extensive dock structure is present over submerged lands at Berths 104, 105, and 106. Historically, Terminal 1 has been used for the staging of lumber, logs, paper products, steel containers, and bagged grain. Various companies have owned or leased portions of the Terminal 1 South Complex (see Remedial Investigation [RI] Report; Hahn and Associates, 2001a). The T1S Site will be redeveloped for residential and commercial purposes. Potentially exposed populations that were evaluated in the HHRA include future residents, current and future commercial workers, and future utility/excavation workers. The site was divided into three Areas of Concern (AOC) and separate risk calculations and risk estimates were conducted for each area. The areas are presented on Figure 2. Human Health Risk Assessment Results for Area A. The assessment of carcinogenic risks to residential receptors at Area A indicated that under both Reasonable Maximum Exposure (RME) and Central Tendency (CT) conditions, the potential risks exceeded DEQ acceptable risk levels. Compounds of Potential Concern (COPCs) that exceeded the Department of Environmental Page 1 Hart Crowser 15191-01-January 18, 2002

	Quality (DEQ) acceptable risk level for individual carcinogens are
	benzo(a)pyrene, benzo(a)anthracene, dibenz(a,h)anthracene,
	benzo(b)fluoranthene, indeno(1,2,3-cd)pyrene and arsenic. The assessment of
•	
:	noncarcinogenic risks identified only lead as present above acceptable risk
	levels for residential exposure under both RME and CT conditions.
	For the commercial worker exposure scenario, the estimated cumulative
	carcinogenic risks were found to be acceptable under both RME and CT
	conditions. However, benzo(a)pyrene and arsenic exceeded the DEQ
	acceptable risk level for individual carcinogens. The assessment of
	noncarcinogenic risks identified lead as present above the acceptable risk level
	for commercial worker exposure under only the RME condition.
 	Tot commission in order of the furthe container.
	For the excavation worker exposure scenario, no unacceptable risks from
	exposure to carcinogens were identified. The assessment of noncarcinogenic
	risks identified lead as present above the acceptable risk level for excavation
	worker exposure under only the RME condition.
	we we are and of only the time contained.
	As discussed in the report, the RME and CT exposure point concentrations
•	(EPCs) for lead in surface and total soil in Area A are driven by the maximum
	detection in one sample (B-68). If this sample were removed from the data set,
	the lead EPCs would be acceptable for the residential and commercial
	receptors. Additionally, while arsenic was identified as a carcinogen resulting in
	unacceptable risks in Area A, there were only two soil samples (within the 0 to
	15 feet depth ranges evaluated in this HHRA) that exceeded the site-specific
	background level of 5.3 mg/kg identified in the RI-(Hahn-and-Associates, 2001a).
·	Human Health Risk Assessment Results for Area B. The assessment of
	carcinogenic risks to residential receptors at Area B indicated that potential risks
	exceeded DEQ acceptable risk level only under the RME condition. COPCs that
	exceed the DEQ acceptable risk level for individual carcinogens are
	benzo(a)pyrene and arsenic. The assessment of noncarcinogenic risks found no
	exceedences of DEQ acceptable risk levels for residential exposure.
	For the commercial worker exposure scenario, the estimated cumulative
	carcinogenic risks were found to be acceptable under both RME and CT
	conditions. However, arsenic exceeded the DEQ acceptable risk level for
•	individual carcinogens under the RME condition. The assessment of
-	noncarcinogenic risks found no exceedences of DEQ acceptable risk levels for
	commercial worker exposure.
	No unaccentable carcinogenic or noncarcinogenic vide was estimated for the
	No unacceptable carcinogenic or noncarcinogenic risks were estimated for the
	excavation worker exposure in Area B.
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_	Arsenic was Identified as a carcinogen resulting in unacceptable risks in Area B
	for residential and commercial worker exposure scenarios. However, there
	were no detected concentrations of arsenic in soils in Area B that exceeded the
	site specific background level of 5.3 mg/kg identified in the RI (Hahn and
	Associates, 2001a).
	Human Health Risk Assessment Results for Area C. The cumulative RME and
	CT carcinogenic risks for all potential receptors (resident, commercial worker,
	and excavation worker) in Area C were found to be acceptable with the
	exception of the RME residential scenario. Arsenic exceeded DEQ individual
	carcinogen-acceptable-risk-level-for the RME residential and commercial
	worker scenarios. The assessment of noncarcinogenic risks found no
	exceedences of DEQ acceptable risk levels for all potential receptors. There
	were no detected concentrations of arsenic in surface soils in Area C that
	exceeded the site specific background level of 5.3 mg/kg identified in the RI
	(Hahn and Associates, 2001a).
	Ecological Risk Assessment Results. The Level 1 Scoping ERA did not identify any
•	ecologically important species or habitats at the T1S Site. The site is almost
	entirely paved or covered by buildings. The absence of upland habitat indicates
, .	that there-are-no-complete exposure pathways for terrestrial ecological-receptors
	to come in contact with contaminated soil at the T1S Site.
	to the military contact may contact the trop bits.
	A Modified Level 2 Screening ERA was conducted on the available groundwater
	monitoring well data collected at this site. There were no detected
	concentrations of organic constituents in the seven groundwater monitoring
	wells that exceeded their corresponding Ecological Screening Benchmark Values
	(SBVs). There were two metals (copper and lead) detected in groundwater that
	exceeded SBVs based on the analysis of unfiltered, total metals, but when the
	same samples were analyzed for dissolved metals, copper and lead were not
	detected. The dissolved fraction of metals represents the bioavailable fraction in
	aqueous environmental media. Therefore, it is concluded that there is no
	potential for adverse ecological impacts to aquatic ecological receptors from the
-	discharge of groundwater to the Willamette River. No additional ecological risk
	assessment activities are warranted at this site.
	assessment activities are warranted at this site.
•	
1.0 IN	TRODUCTION AND PURPOSE
	· · · · · · · · · · · · · · · · · · ·
-	This report summarizes the results of the human health risk assessment (HHRA)
	and Level 1 Scoping and Modified Level 2 Screening ecological risk assessment
	(ERA) performed at and in the vicinity of the Port of Portland (Port) Terminal 1
Hart Crows	er Page 3
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South (T1S Site) in Portland, Oregon. This report was prepared for the Port, Project Number 24232 and Task Number 730. The purpose of the HHRA is to evaluate current and predicted future site conditions and to assess if these conditions pose unacceptable risks to public health. The purpose of the Level 1 Scoping ERA is to provide a conservative, qualitative determination of whether ecological receptors and/or exposure pathways are potentially present at or inthe locality of the site. The Modified Level 2 Screening ERA was conducted on site groundwater data to determine whether constituents were present at levels of concern for aquatic ecological receptors. The HHRA was conducted in accordance with the protocol for performing risk assessments-under-Oregon-Administrative-Rules (OAR)-340-122-084 and the Department of Environmental Quality's (DEQ's) Guidance for Conduct of Deterministic Human Health Risk Assessments (DEQ, 2000). Additionally, the scope of this risk assessment was further defined based on the Risk Assessment Work Plan (Hart Crowser, 2001), DEQ-Comments on the Risk Assessment-Work Plan (letter dated October 25, 2001), and Port of Portland's Response to Review Comments (letter dated November 12, 2001). The Level 1 - Scoping ERA was completed in accordance with the Guidance for Ecological Risk Assessment (DEQ, 1998) and the Modified Level 2 Screening ERA was completed in accordance with the methodology presented in the Risk Assessment Work Plan and further discussed with DEQ in the Port of Portland's Response to Review Comments Letter. This report is organized as follows: Section 2.0 - Site Location and History Section 3.0 - Human Health Risk Assessment Section 4.0 - Level 1 Scoping Ecological Risk Assessment Section 5.0 - Limitations Section 6.0 - References 2.0 SITE LOCATION AND HISTORY This section summarizes the available information on this site. A more detailed description-of-environmental-activities-and-the-results-of-the-remedial investigation (RI) conducted at this site are provided in the Terminal 1 South Remedial Investigation Report (Volumes 1 and 2) prepared by Hahn and Associates (Hahn and Associates, 2001a). Hart Crowser Page 4 15191-01-January 18, 2002

2.1.1 Site Location The T1S Site is located at 2100 NW Front Avenue along the Willamette River in Portland, Oregon (Figure 1). The site consists of approximately 21 acres located
2.1.1 Site Location The T1S Site is located at 2100 NW Front Avenue along the Willamette River in
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The T1S Site is located at 2100 NW Front Avenue along the Willamette River in
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Portland, Oregon (Figure 1). The site consists of approximately 21 acros located
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northwest of Interstate 405 (Fremont Bridge), northeast of NW Front Avenue,
southeast of Slip No. 2, and southwest of the Willamette River (Figures 1 and 2).
The T1S Site does not include sediments adjacent to the Site.
2.1.2 Site Description
Two primary structures, designated as Warehouse No. 2 and House No. 104,
are currently located at the T1S Site. Tristar Transload currently leases and
operates the open storage area southeast of Slip No. 2 and northwest of House
No. 104 and portions of House No. 104. The remaining portions of the site are
unoccupied. Additionally, an extensive dock structure is present over
submerged land at Berths 104, 105, and 106.
The topography at the T1S Site is generally level at an elevation of
approximately 30 feet above mean sea level (msl). The site is generally paved
with asphalt or concrete with little or no vegetation or bare ground present.
2.1.3 Site History
The site history presented here is summarized from information contained in a
Preliminary Assessment (PA) (Port of Portland, 2000) prepared for the T1S Site.
In approximately 1884, upland areas in the vicinity of Terminal 1 extended 100
to-200-feet-northeast of Front Avenue. By 1908, they extended approximately—
200 to 400 feet northeast of NW Front Avenue. Since that time, various
portions of the T1S Site have been filled and dredged. Slip Nos. 1 and 2 were
created by dredging in approximately 1914 and 1923, respectively. Filling
activities at the site were generally completed in approximately 1972 when Slip No. 1 was filled.
Between 1913 and 1936, the Commission of Public Docks purchased various
parcels of property in four primary phases. Three of these parcels now make up
the Marine Terminal 1 South complex. The Commission of Public-Docks
merged into the Port on January 1, 1971.
Prior to and during World Was II Tamainal 1 and 11 and 11 and 1
Prior to and during World War II, Terminal 1 and the adjacent industrial neighborhood supported expanded activities on behalf of the war effort. Ship
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	building and repair at the Willamette Iron and Steel facility formerly located at
·	Terminal 1 necessitated increased dock front dredging (for larger ship berths)
•	and the occasional use of Terminal 1-property for temporary equipment storage.
	In 1946, the Commission of Public Docks (CPD) purchased the Eastern and
	Western Lumber Company property to the immediate north of Terminal 1
	South. Willamette Iron & Steel Corporation, now adjacent to the CPD
	terminal, changed ownership in the same year, becoming the Willamette Iron
	and Steel Company.
	Historically, Terminal 1 has been used for the staging of lumber, logs, paper
	products, steel containers, and bagged grain. Various companies have owned
	or leased portions of the Terminal 1 South Complex (see RI Report; Hahn and
•	Associates, 2001a). The T1S Site will be redeveloped for residential and
	commercial purposes.
2.2 Site (Geology and Hydrogeology
	This section presents a summary of the site geology and hydrogeology.
	Additional details of site geology and hydrogeology are presented in the RI
	Report (Hahn and Associates, 2001a) and the Monitoring Well Installation and
<u> </u>	Groundwater Sampling Report (Hahn and Associates, 2001b).
<u>.</u>	2.2.4 Coology
	2.2.1 Geology
	The subsurface soils encountered during the investigations were
	predominantly sands and silts with occasional gravel to the maximum depth
	of investigation at 80 feet below ground surface (bgs).
	■ Based on historical documentation and investigations, the property has been
	extensively filled in through time; fill material was encountered at all push-
	probe locations from the surface to depths of 32 to 67 feet bgs.
	Soils thought to be former Willamette River sediments were encountered at
	the former Slip No. 1 (B-84) at a depth of approximately 67 feet bgs.
· · ····	Soils encountered beneath NW Front Avenue were generally siltier than
	those encountered on the T1S Site, suggesting the soils in the right of way
	are either alluvial in origin or from a different fill source than that of the site.
	2.2.2 Hydrogeology
	■ Groundwater in the vicinity of the T1S Site generally occurs in three
	principal hydrogeologic zones: (1) a shallow unconfined fill/alluvial deposit
·	(shallow water-bearing zone [WBZ]); (2) generally confined Troutdale WBZ;
	(Similar state) Searn & Folio (11921), (2) Benefally Committee Hodicale WDZ,
	Page 6

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	and (3) the confined Columbia River Basalt WBZ.
	 Unconfined groundwater was encountered within the shallow WBZ (fill) at
	an average depth of approximately 23 feet bgs.
	■ Groundwater elevation measured in the seven monitoring wells installed at
	the T1S Site on September 28 and October 30, 2001 indicate a general flow
	to the northeast towards the Willamette River with a decline or even reversal
·	of the gradient near the river (Hahn and Associates, 2001b).
2.3 Previo	us Site Investigations
2.5 Fievio	us Site investigations
	In July 2001, Hahn-and-Associates-completed an-RI at the T1S-Site (Hahn-and
	Associates, 2001a). RI activities completed at this site consisted of the following
	five phases:
	■ Focused Environmental Site Assessment-completed by Maul Foster in 1998
•	(Maul Foster, 1998);
	■ Environmental Baseline Investigation completed by Hahn-and-Associates in
	February and March, 2000 (Hahn and Associates 2001a);
	■ B-38 Area Characterization completed by Hahn and Associates in March
	2000 (Hahn and Associates 2001a);
	■ Supplemental Site Characterization Activities completed by Hahn and
	Associates in September 2000 (Hahn and Associates 2001a); and
	 Data Gap Investigation completed by Hahn and Associates during October and November 2000 (Hahn and Associates 2001a).
	and November 2000 (Hann and Associates 2001a).
	A total of 112 push-probe borings were installed for the collection of soil and
	groundwater samples during these site activities. The locations of these push-
	probe borings are presented on Figure 2. Please refer to the RI Report (Hahn
	and Associates, 2001a) for further discussion of these activities and results.
	A groundwater investigation was conducted at the T1S Site by Hahn and
	Associates in August, September, and October 2001 (Hahn and Associates
	2001b). Site activities included installation, development, and sampling of seven groundwater monitoring wells at the site. The locations of the groundwater
•	monitoring wells are presented on Figure 2. Please refer to the groundwater
	Associates, 2001b).
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2.4 Beneficial Land and Water Use Surveys

Beneficial land and water use determinations were completed at the T1S-Site-to-identify current and reasonably likely future uses of land and water in the vicinity of the Site. This information was presented in the RI Report (Hahn and Associates, 2001a) and used to ensure that appropriate exposure scenarios were selected for evaluation in the proposed RA.

2.4.1 Locality of the Facility

The locality of the facility (LOF) is defined as "any point where a human or ecological receptor contacts, or is reasonably likely to come into contact with, facility related hazardous substances."

Chemicals have been detected in both soil and groundwater at various areas of the site, but off-site migration of contamination is not evident based on the existing data. Accordingly, the LOF is defined only as the T1S Site and the adjacent area on Front Avenue in Area A (Figure 2).

2.4.2 Land Use

Historical Land Use. The approximate 21-acre T1S Site has historically been zoned as "IH" for Heavy Industrial. Surrounding adjacent properties are zoned "IH" Heavy Industrial and "EX" Central Employment.

Current and Reasonably Likely Future Land Use. The current and reasonably likely future land use in the LOF is well defined. The site is currently zoned as Central Residential (RX) such that it can be redeveloped for an alternative use. The RX zoning is considered the comprehensive plan for the property. Based on the RX zoning designation, it is expected the site will be used for mixed-use residential/commercial development in the future.

2.4.3 Groundwater Beneficial Use

A beneficial groundwater use evaluation was conducted for the Hoyt Street Property (RETEC, 1997) that adjoins the southeast corner of the T1S Site. Flahn and Associates conducted an additional well inventory as part of the RI and the groundwater monitoring study to supplement the RETEC survey. Based on trends in groundwater use in the area as well as RETEC fate and transport modeling, the only identified beneficial use for groundwater in the LOF is discharge to the Willamette River. No water wells were found to be in use within one-half mile of the T1S Site. No surface water rights were identified within one-half mile of the T1S Site.

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2.5 Chemical Data Quality Review Prior-to-identifying Compounds of Potential Concern (COPCs) for the T-IS-Site; a chemical data quality review was conducted on the available soil and water analytical data collected as part of the RI completed at this site (Hahn and Associates, 2001a and 2001b). The following criteria were evaluated in the data quality review process: Holding times; Method blanks; -Surrogate-recoveries: Laboratory control sample/laboratory control sample duplicate (LCS/LCSD) recoveries; Matrix-spike/matrix-spike duplicate (MS/MSD)-recoveries; and Laboratory and field duplicate relative percent difference (RPD). The results of this data quality review are presented in Appendix F. Only those data of sufficient quality for use in the risk assessment were carried forward for COPC screening and risk calculations. The data that did not meet data quality objectives were rejected because quality assurance samples were not runconcurrently with site samples. The exclusion of this data had no effect on the objectives of this risk assessment. The data that were rejected based on data quality concerns are discussed below: Diesel and Oil. Nine diesel/oil sample results from the Area A data set were rejected. However, only five of these samples were from the depth profile of 0 to 15 feet bgs that was considered in the risk assessment. Fifty-three diesel/oil samples were included in the Area A risk assessment data set. Eight diesel/oil sample results from the Area B data set were rejected (six in surface soil and two-in-subsurface soil), while 30 diesel/oil samples were included in the Area B risk assessment data set. All diesel and oil sample results from the Area B rejected samples were either low level or nondetect. BTEX. Six BTEX sample results from the Area A data set were rejected based on data quality concerns. All of the BTEX results from the rejected samples and from the samples that were not rejected were nondetect. In addition, all VOC samples collected at Area A-were nondetect for BTEX. PAHs. Two PAH sample results from the Area A data set were rejected (one within the 0 to 15 feet depth profile), while 41 PAH samples were included in the risk assessment data set. Hart Crowser Page 9 15191-01 January 18, 2002

	■ PCBs. One PCB sample result from the Area A data set was rejected based
	on data quality concerns (sample-B-38 collected at 10 feet bgs). The PCB
	sample results for this sample were nondetect.
2.6 Identific	ation of Compounds of Potential Concern
	Chemical analyses on samples collected at the T1S Site have identified diesel
	and oil as the fuel types present. However, due to the current lack of toxicity
	data for diesel or oil as a whole (each fuel type is a complex mixture of hundreds
	of chemical compounds), these fuels were not quantitatively evaluated in the
	HHRA. Instead, we focused on individual petroleum constituents within these
	fuel types-for which appropriate-toxicity data-are available.
	Total type State will appropriate toxicity data are available.
	Specific chemical constituents of these fuel types are possible compounds of
	interest (COI). COIs are defined as compounds detected at the site, and COPCs
	are those COIs that exceed the risk-based screening levels as discussed below
•	and are carried forward in the HHRA. Based on investigations conducted at the
	T1S Site, the COIs in soil and groundwater include the following groups of
	compounds: Total Petroleum Hydrocarbons (TPH), polycyclic aromatic
	hydrocarbons (PAHs), volatile organic compounds (VOCs), polychlorinated
	biphenyls (PCBs), and metals.
	Evaluation for COPCs. In accordance with DEQ human health risk assessment
	guidance (DEQ, 2000), soil COIs were conservatively screened against EPA
	Region 9 Residential Soil Preliminary Remediation Goals (PRGs), and
	groundwater COIs were conservatively screened against EPA Region 9 Tap
•	Water PRGs (EPA, 2000a). Because exposure to groundwater is limited to
	inhalation of VOCs that have migrated from groundwater to indoor or outdoor
<u>. </u>	air, only VOCs detected in groundwater were evaluated as potential COPCs.
-	`
	Additional steps, which are described in Section 2.3.2, (3)(a) through (e) of the
	DEQ human health risk assessment guidance, were also performed to ensure
	potential cumulative effects from multiple compounds or from an individual
	compound detected in multiple media were accounted for.
	As presented in the Risk-Assessment-Work Plan for this-site (Hart Crowser,
	2001), the site was divided into three separate Areas of Concern (AOC). The
	AOCs are presented in Figure 2. COPCs were identified for each area and
	separate risk calculations and risk estimates were conducted for each area.
	, and the row community were conjudeted for each area.
	Tables 1 through 3 summarize the identification of COPCs in soil and
	groundwater for Areas A, B, and C, respectively. The following COPCs were
F	identified in each area:
<u> </u>	

Area A

- Soil: Diesel, oil, benzo(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, dibenz(a,h)anthracene, indeno(1,2,3-cd)pyrene, arsenic, and lead. No soil PRGs are available for diesel and heavy oil; however, since both analytes were detected in soil they were retained as COPCs.
- Groundwater: Diesel and tetrachloroethene. Diesel was identified as a COPC because a tap water PRG is not available. Heavy oil was not detected in groundwater.

Area B

- Soil: Diesel, oil, benzo(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, dibenz(a,h)anthracene, indeno(1,2,3-cd)pyrene, and arsenic. Diesel and oil were identified as COPCs because soil PRGs are not available.
- Groundwater: Diesel. Diesel was identified as a COPC because a tap water PRG is not available. Heavy oil was not detected in groundwater. Tetrachloroethene was not identified as a COPC for Area B since it was only detected in Monitoring Well 1, which is located in Area A.

Area C

- Soil: Arsenic.
- Groundwater: None.

3.0 HUMAN HEALTH RISK ASSESSMENT

This risk assessment conforms to the protocol for performing risk assessments under OAR 340-122-084 and DEQ's Guidance for Conduct of Deterministic Human Health Risk Assessments (DEQ, 2000). Other guidance were used as appropriate and where indicated. The HHRA evaluates the probability and magnitude of adverse impacts on human health associated with actual or potential exposure to site-related COPCs. This information was used to determine what additional remedial actions are needed (if any) to mitigate any predicted impacts. Deterministic human health risk assessments for both existing and reasonably likely future exposure scenarios were performed.

In accordance with EPA and DEQ guidance, the risk assessment consists of the

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following four phases: Exposure Assessment, Toxicity Assessment, Risk Characterization, and Uncertainty Analysis. In the exposure assessment, potentially exposed populations and potentially complete exposure pathways (shown in the human health conceptual site model [CSM], Figure 3) were identified based on current and future land use scenarios. Exposure point concentrations (EPC) and reasonable maximum exposure (RME) and central tendency (CT) intake rates were calculated for each complete exposure pathway based on the use of exposure factors that reflect site-specific conditions. In the toxicity assessment, quantitative toxicity information was collected, and appropriate toxicity values were determined for use in quantifying carcinogenic and non-carcinogenic risks associated with exposure to site-related chemicals In the risk characterization phase, the results of the exposure assessment and toxicity assessment were combined to estimate the potential cancer risks and non-cancer hazard quotients at the site. In the uncertainty section, the uncertainty associated with the exposure assessment, toxicity assessment, and risk characterization sections are discussed. 3.1 Exposure Assessment The objectives of an exposure assessment are to: Identify potentially exposed populations; Identify potentially complete exposure pathways; and Measure or estimate the magnitude, duration, and frequency of exposure for each receptor (or receptor group). 3.1.2 Final Conceptual Site Model The final conceptual site model (CSM) is based on an evaluation of existing data and the current and reasonably likely future conditions at the site (Figure 3). This model provides the framework for assessing potential exposure pathways to be considered in the risk assessments. To be considered complete, an exposure pathway must have: (1) an identified source of COPCs; (2) a release/transport mechanism from the source; and (3) a receptor to which contact can occur. At this site, likely or potential sources include former USTs, former ASTs, machine shop areas, paint/battery/waste oil/drum/chemical storage, railroad spur, and miscellaneous spills and leaks. Potentially Exposed Populations. A beneficial land and water use survey has

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been completed for the site and is discussed in Section 2.4 (Hahn and Associates, 2001). Based on the Central Residential (RX) zoning designation, it is expected that the site will be used for mixed-use residential/commercial development in the future. The only identified beneficial use for groundwater in the locality of the facility is discharge to the Willamette River. Therefore, the final CSM assumes the future area land use will be a mix of residential and commercial and that groundwater beneath the site is not and likely will not be used for drinking water. Figure 3 presents the final CSM for this site. The red boxes on the figure indicate potentially complete pathways to the indicated receptor. In addition to residential and commercial receptors, the HHRA-will-also-evaluate utility/excavation workers as potentially exposed populations. Utility/excavation workers will be identified as excavation workers in the remainder of the HHRA. Potentially Complete Exposure Routes. Exposure pathways for quantitative analysis were selected based on the final CSM developed for this site. Based on available information, the exposure pathways evaluated in this HHRA are: Incidental ingestion of soil (all receptors); Dermal contact with soil (all receptors): Inhalation of fugitive dust from surface soil (residents and commercial workers): Inhalation of fugitive dust from total soil (0 to 15 feet below ground surface, excavation workers); and Inhalation of VOCs from groundwater (all receptors; indoor for residents and commercial workers; outdoor air only for excavation workers). No VOCs were identified as soil COPCs (see Tables 1 through 3). Outdoor air was not evaluated for residents and commercial workers since the risks and hazards associated with indoor air, which are higher than those associated with outdoor air, were acceptable. Direct contact with groundwater is not considered a potential exposure pathway for excavation workers, as the average depth of the shallow WBZ was reported to be 23 feet bgs (Hahn and Associates, 2001a). For this HHRA, residents and commercial workers are assumed to be exposed to soil down to a depth of 3 feet below ground surface (bgs) and excavation workers are assumed to be exposed to-soils-down-to-a-depth-of-15-feet-bgs-Areas of Concern. The T1S Site is being redeveloped for residential and commercial purposes. The site will be developed into three areas (A, B, and C) Hart Crowser Page 13 15191-01-January-18, 2002

which were evaluated as separate areas of concern (AOCs). Separate COPCs were identified and separate risk calculations conducted for each AOC. The RI identified six general areas/locations of soil impacted with petroleum hydrocarbons. Area A includes the B-20 Area, B-38 Area, and B-102 Area. Area B includes the B-5 Area, B-29 Area, and B-37 (Dry Well) Area. Area C does not include any areas/locations of soil impacted with petroleum hydrocarbons. The AOCs for this site are presented on Figure 2. 3.1.2 Development of Exposure Point Concentrations Exposure point concentrations (EPCs) represent the chemical concentrations in the soil and groundwater that the receptor-will potentially contact during the exposure period. The EPCs for the site's COPCs were derived from either data obtained from sampling or from a combination of sample data and fate and transport modeling. For example, air EPCs were modeled from groundwater EPCs for volatile constituents. Groundwater data from monitoring well samples collected in September and October 2001 were used to represent current and future groundwater conditions. The residential and commercial worker scenarios were evaluated based on exposure to surface soil (0 to 3 feet bgs), while the excavation worker scenario was based on exposure to surface and subsurface soil (0 to 15 feet bgs). No VOCs were Identified as soil COPCs, therefore, soil from 15 feet bgs down to groundwater was not considered in the volatilization to indoor and outdoor air pathways. In accordance with EPA guidance (EPA, 1989) for chemicals detected at one sampling location but not at others, a proxy concentration equal to half the sample quantification limit (SQL) were used to represent the concentration of the chemical of concern in a sample where it is not detected. The 90 percent upper confidence limit (UCL) on the arithmetic mean concentration of COPCs in each environmental medium of concern were used to evaluate the reasonable maximum exposure (RME) scenario, while the arithmetic mean were used to evaluate the central tendency (CT) exposure scenario (EPA, 1989). The RME scenario is intended to be a conservative estimate of potential exposure, while the CT exposure scenario is intended to be a more realistic exposure scenario. Using both the RME and CT allows for a range of potential risk and hazard estimates. The 90 percent UCL is calculated based on EPA (1992) guidance. The manner of calculating the 90 percent UCL were as follows: As a first step, the underlying distribution of the data was evaluated using the Hart Crowser Page 14 15191-01 January 18, 2002

Shapiro and Wilk W-Test (Gilbert, 1987) to determine if the data are normal or lognormal. If the normal and lognormal distributions are indicated, the 90percent UCL were calculated appropriately. If the normality test rejects both normal and lognormal distributions at a significance level of 95 percent, the test was rerun by adjusting the W-Test quantile downward by 0.1 from the original quantile (providing a greater tolerance for accepting a distribution). If the data set conforms to a normal or-lognormal distribution with the greater-tolerance, the distribution was reported as weak lognormal (or weak normal). If the normal and lognormal distributions are rejected with the greater tolerance, the data were assumed to fit a lognormal distribution for calculation of the 90 percent UCL (assumed lognormal distribution; EPA, 1992). In cases where the 90 percent UCL or the calculated mean concentration exceed the maximum detected value (which can occur in data sets with a large variance), the maximum detected value were used to define the upper limit of this range. EPCs for this HHRA are presented in Table 4. All of the EPCs presented in Table 4 were calculated using the methodology presented above, with the following exceptions: Area A: Total Soil (0 to 15 feet bgs). The 90 percent UCL for benzo(a)anthracene of 0.35 mg/kg is less than the arithmetic mean of 0.37 mg/kg. This is primarily due to the elevated detection of 9.35 mg/kg, which was detected in the soil sample B-68. The RME concentration for benzo(a)anthracene was, therefore, set at 0.37 mg/kg. Area B: Total Soil (0 to 15 feet bgs). The 90 percent UCLs and arithmetic means for the cPAHs were significantly affected by the elevated SQL of 67 mg/kg in soil sample B-63. Since one half of 67 mg/kg is over 10 times greater than the maximum detected concentrations of the five cPAHs, the cPAH results for soil sample B-63 were not included in the statistical evaluation for this data set. However, subsurface soil samples collected adjacent to sample B-63 (e.g., Samples B-66 and B-67) had detected concentrations of cPAHs that were included in the calculations of EPCs. Therefore, this area was represented in the calculated risk estimates and the exclusion of cPAH SQLs from Sample B-63 had no impact on HHRA objectives. The inhalation of particulates and VOCs pathways were evaluated using the fate and transport models presented in DEQ's risk assessment guidance (DEQ, 2000) and risk-based decision-making guidance (DEQ, 1999).

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3.1.3 Exposure Factors To quantitate intake estimates for site-related chemicals, EPCs are combined with variables that describe the exposed population (e.g., contact rate, exposure frequency and duration, body weight). Exposure factors were selected using standard default exposure factors presented in Guidance for Conduct of Deterministic Human Health Risk Assessments (DEQ, 2000). Industrial exposure assumptions were used to evaluate the commercial scenario. The following paragraphs describe the exposure pathways proposed for evaluation in this HHRA. Incidental Soil Ingestion. Incidental ingestion of soil is often a primary route of exposure to particulate-bound chemicals. Individuals have been observed to ingest small amounts of soil as a result of hand-to-mouth behavioral patterns that may follow soil contact activities. RME and CT factors applicable to this pathway for the identified human receptors are summarized in Table 5. Dermal Soil Contact and Absorption. In addition to leading to incidental soil ingestion, soil contact can also result in absorption of some chemicals directly through the skin. RME and CT exposure factors for the dermal contact pathways are summarized in Table 6. Dermal absorption rates have not been well defined in the available literature. Current RME and CT dermal absorption factors were selected from DEQ Human Health Risk Assessment Guidance (DEQ, 2000). Air Inhalation. Exposure to chemicals present in soil and groundwater may also result from inhalation of vapors and/or fugitive dust generated at the site. RME and CT factors applicable to this pathway are summarized in Tables 7 and 8. 3.2 Toxicity Assessment The objectives of the toxicity assessment are to evaluate the inherent toxicity of the compounds under investigation and to identify and select toxicological measures for use in evaluating the significance of the exposure. These toxicological measures or criteria were used in conjunction with intake rates for chemicals of concern in the risk characterization process of the human health risk assessment. Standard human health risk assessment toxicity databases were used to derive health-based toxicity criteria. The hierarchy of sources for toxicity criteria for use in this risk assessment will follow as presented in OAR 340-122-084. The hierarchy of toxicity criteria is as follows:

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(1) EPA's Integrated Risk Information System (IRIS, EPA 2000b); (2) EPA's Health Effects Assessment Summary Table (HEAST, EPA 1997); (3) EPA-NCEA Superfund Health Risk Technical Support Center; (4) Other U.S. EPA documents or databases; (5) ATSDR minimal risk levels (MRLs); and (6) Other professionally peer-reviewed documents as needed and as approved by DEQ. 3.2.1 Types of Toxicity Values for Quantifying Risks Toxicity and risk assessments vary for different chemicals depending upon whether non-carcinogenic or carcinogenic responses (i.e., endpoints) are used to assess potential risks. These criteria, in turn, are based on the endpoints observed from laboratory or epidemiological studies with the chemicals. Some chemicals of concern may result in both non-carcinogenic and carcinogenic effects, although in many cases the EPA has published toxicity criteria for only the most sensitive type: of toxic effect supporting the most restrictive toxicological criteria. The toxicity criteria used in this HHRA are presented in Table 9. Reference Doses (RfDs). Reference doses are used to quantitatively evaluate non-carcinogenic toxicity of a specific chemical. Reference doses are established at levels associated with no adverse effect—the "no observed adverse effect level" (NOAEL). In general, the RfD is an estimate (with uncertainty spanning perhaps an order of magnitude) of a daily exposure to the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a lifetime. RfDs are developed from an analysis of the available toxicological literature from which a critical study is selected. The selection of a critical study is made by professional judgment and considers factors such as the quality of the study, the relevance of the study to human exposures, and other factors. Good quality human toxicological data are preferred to animal studies. If human data are not available, the study on the most sensitive species is selected as the critical study. Similarly, the toxic effect manifested at the lowest exposure level is (generally) selected as the critical effect. Cancer Slope Factors (SFs). The toxicity of potential human carcinogens is evaluated differently. It is assumed for carcinogens that no threshold concentrations exist below which adverse effects may not occur. Probabilistic methods based on chemical-specific dose-response curves are used to establish slope factors, which are Hart Crowser Page 17 15191-01 January 18, 2002

then used to quantify potential risks from exposure to carcinogens.

Dose-response curves are generated in laboratory studies using high chemical concentrations. The dose-response curve is fitted to a linearized multistage model that extrapolates the slope of the curve from high experimental concentrations to low concentrations at which people are typically exposed. The final slope factor (SF) is based on the 95 percent UCL of the extrapolated slope of the dose-response curve. Because of the non-threshold assumption and the UCL statistical procedure, the use of published slope factors provides a conservative upper-bound estimate of potential risks associated with exposure.

3.2.2 Modification of Oral Toxicity Values for Evaluating Dermal Exposure

Oral toxicity values are expressed as administered doses. When evaluating dermal exposure to contaminants from soil and water, it is necessary to adjust the oral toxicity value (which is based on an administered dose) to one based on an absorbed dose using a chemical's oral absorption efficiency. However, according to EPA guidance (2000a), the only chemical for which an adjustment is recommended at this time is cadmium. Adjustment is not recommended for other chemicals because a scientifically defensible database does not exist for making the adjustment. Therefore, in this HHRA, because cadmium is not a COI at this site, no adjustments of oral toxicity factors to evaluate dermal exposure were done.

3.2.3 Toxicity Assessment for Lead

Lead is a unique chemical in its pharmacokinetic and toxicological properties. Although classified as both a potential carcinogen (B2 weight of evidence) and a non-carcinogen, lead is most often assessed as a non-carcinogen only, since these effects manifest themselves at doses lower than those for carcinogenicity. However, in contrast to the assumption of the existence of a threshold for non-carcinogenic responses, there does not appear to be a threshold below which lead does not exert a response.

Currently, the EPA provides neither a reference dose for evaluating the noncarcinogenic effects (unrelated to cancer) nor a slope factor for evaluating the carcinogenic effects for lead. EPA has developed an exposure model for lead that considers both its biokinetics and toxicological properties. This model—the "Integrated Exposure Uptake and Biokinetic" (IEUBK) model—integrates the intake of lead from multiple sources, including soil, food, and water ingestion, inhalation, and, when appropriate, maternal contributions. Intakes are assessed for children from ages 0 (birth) to seven. The model does not assess lead

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intakes for older children or adults. Childhood exposure to lead is the focus of this model because this receptor group is recognized as the most sensitive to the non-carcinogenic effects of inorganic lead. Therefore, to evaluate lead exposures at the T1S Site, we will use other criteria as described below. Soil exposures at the site are limited to residents and commercial workers contacting soil at the ground surface or excavation workers contacting soil during trenching or excavation activities. We screened the soil lead concentrations against the EPA Region 9 residential soil PRG for lead (400 mg/kg) to evaluate residential exposure and the adult soil screening level for lead (750 mg/kg) to evaluate commercial and excavation exposures. Lead has been detected in groundwater at the site. However, as lead is not volatile and no direct contact exposure pathways have been identified to human receptors, lead in groundwater will not be further evaluated in the HHRA. 3.2.4 Toxicity Assessment for Total Petroleum Hydrocarbons (TPH) Determining appropriate toxicity values for TPH (a class of compounds identified as a COI at this site) is difficult because of the characteristics of TPH. TPH are a complex mixture of hundreds or more individual alkanes, cycloalkanes, alkenes, aromatics, and other petroleum substances. For this HHRA, the human health risks associated with TPH were evaluated using an indicator approach. The indicators refer to single compounds within TPH known or believed to be carcinogenic and non-carcinogenic and which are evaluated individually. The indicator compounds that were quantitatively evaluated in this HHRA are: Volatiles (BTEX): benzene, toluene, ethylbenzene, and xylene; and Polynuclear Aromatic Hydrocarbons (PAHs): anthracene, acenaphthene, benzo(a)pyrene, pyrene, naphthalene, chrysene, ideno(1,2,3-cd)pyrene, benzo(k)flouranthene, flourene, naphthalene, benzo(b)flouranthene, benzo(a)anthracene, and dibenzo(a,h)anthracene. 3.3 Risk Characterization Risk characterization is the process of comparing the chemical intake by a receptor to the toxicity of the chemical. This comparison is expressed either as a hazard index-(non-carcinogens)-or-an-excess-lifetime risk-of cancer (carcinogens). Hart-Crowser Page 19 15191-01 January 18, 2002

	3.3.1 Methods Used to Quantify Risks and Hazards	
	The monitors used to Quantity Risks and Hazarus	•
	As discussed in Section 3.2, non-carcinogenic chemical effects are	
	quantitatively evaluated using a RfD, while carcinogenic chemical effects are	
	evaluated using a SF.	
		•
	Non-Carcinogenic Effects. For the residential exposure pathway, the non-	
	carcinogenic intakes are based on child exposures, which are more conservative	
	than potential adult exposures. The daily intake of each compound resulting	· · · · · ·
	from site exposure is divided by the available RfD value for the compound to	
	compute a hazard quotient (HQ), as follows:	
· .		
 	HQ = CDI/RfD	
	where:	
	· · · · · · · · · · · · · · · · · · ·	
	CDI = Chronic daily intake; the estimated exposure level over a given time	
	period in mg/kg-day.	
	RfD = Reference Dose; the exposure level that is likely to be without	
•	deleterious effects during a given time increment in mg/kg-day.	
	Only-chronic RfDs were used for this risk assessment.	<u>····</u>
	Consinguity PSS at Fourth 11 at 1	
	Carcinogenic Effects. For the residential exposure pathway, the carcinogenic	
	intakes are based on combined adult and child exposures, which are more	
	conservative than child or adult exposures calculated separately. An estimated	•
	excess lifetime cancer risk is calculated using:	
•	Risk = CDI x SF	
	NISK — CLAI X SF	
	where:	
	MIICIC'	
	CDI = Chronic daily intake; the estimated lifetime exposure level in	
•	mg/kg-day.	
	mg/kg-uay.	
	SF = Slope-Factor; the upper-bound estimate of the probability of a	
	cancer response per unit of intake of a chemical over a lifetime,	
	expressed as (mg/kg-day) ⁻¹ .	
	entrepos de finition de la constantina della con	
	Cumulative Hazard and Risk Estimates. For simultaneous exposure to multiple	•
	chemicals with similar toxic effects or target organ, a Hazard Index (HI) is	•
· · · · · ·	calculated as the sum of chemical-specific HQs. A toxic effect is considered	.
	possible if a HI or HQ exceeds 1 (OAR 340-122-115).	
	possession a fit of Fig. coccoods 1 (O/AC 370-122-113).	٠
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For simultaneous exposure to multiple chemicals, individual risk estimates are summed to provide pathway, media, and receptor total risk estimates.

Combining potential cancer risks as a result of exposure to multiple chemicals through multiple exposure pathways assumes the following:

- Exposure to all COPCs will result in the same effect (cancer); and
- Each COPC exerts-its-effect independently (i.e., there is no synergismor antagonism).

OAR 340-122-115 considers 1 x 10⁻⁶ and 1 x 10⁻⁵ to be acceptable risk levels for individual and multiple-carcinogens, respectively.

3.3.2 Risks and Hazards Associated with Current and Future Site Conditions

Risk and hazard estimates for each area are discussed in Sections 3.3.2.1 through 3.3.2.3. As discussed previously, the residential, commercial worker, and excavation worker exposure scenarios are evaluated for each area. Risk and hazard estimate calculations for each area, exposure pathway, and receptor are presented in Appendix B (Tables B-1 through B-9). Table 10 presents the totals as a sum of risks and hazards associated with each individual exposure pathway, while Table 11 presents the RME carcinogenic risk estimates as a sum of risks associates with each COPC.

3.3.2.1 Area A Risk and Hazard Estimates

The exposure pathways that were quantitatively evaluated at Area A are soil ingestion, dermal contact with soil, inhalation of volatiles from groundwater, and inhalation of fugitive dust.

Resident. The cumulative RME and CT excess lifetime cancer risks for the residential receptor are estimated to be 2×10^4 and 2×10^6 , respectively. The RME risk estimate is greater than the OAR 340-122 acceptable level of 1×10^5 for cumulative carcinogenic risk, while the CT risk estimate is less than the DEQ acceptable risk level. The primary exposure pathways (RME evaluation) are soil ingestion (risk = 8×10^5) and dermal contact with soil (risk = 8×10^5). COPCs that exceed the DEQ acceptable risk level of 1×10^6 for individual carcinogens are benzo(a)pyrene (risk = 9×10^5), arsenic (risk = 5×10^6), benzo(a)anthracene (risk = 1×10^5), dibenz(a,h)anthracene (risk = 8×10^6), benzo(b)fluoranthene (risk = 7×10^6), and indeno(1,2,3-cd)pyrene (risk = 4×10^6).

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The cumulative RME and CT HIs for the residential receptor are estimated to be 1.0 and 0.01, respectively: The RME HI is equal to, while the CT HI is less than, the DEQ acceptable HI of 1.0. Commercial Worker. The cumulative RME and CT excess lifetime cancer risks for the commercial worker are estimated to be 1-x-10⁻⁵ and 6 x 10⁻⁷, respectively. The RME risk estimate is equal to, while the CT risk estimate is less than, the DEQ acceptable level of 1 x 10⁻⁵ for cumulative carcinogenic risk. The primary exposure pathways (RME evaluation) are soil ingestion (risk = 1×10^{-5}) and dermal contact with soil (risk = 3×10^6). Individual COPCs that exceed the DEQ acceptable risk level of 1 x 10⁶ for individual carcinogens are benzo(a)pyrene (risk = 7×10^6) and arsenic (risk = 5×10^6). The cumulative RME and CT HIs for the residential receptor are estimated to be 0.03 and 0.006, respectively. The RME and CT HI are less than the DEQ acceptable HI of 1.0. Excavation Worker. The cumulative RME and CT excess lifetime cancer risks for the excavation worker are estimated to be 5×10^8 and 3×10^9 , respectively. The RME and CT risk estimates are less than the DEQ acceptable risk level for multiple carcinogens. The cumulative RME and CT HIs for the residential receptor are estimated to be 4 x 10³ and 5 x 10⁴, respectively. Both of which are well below the acceptable HI of 1.0. 3.3.2.2 Area B Risk and Hazard Estimates The exposure pathways that were quantitatively evaluated at Area B are soil ingestion, dermal contact with soil, and inhalation of fugitive dust. No VOCs were detected in Area B-soil or groundwater. Resident. The cumulative RME and CT excess lifetime cancer risks for the residential receptor are estimated to be 3×10^{-5} and 5×10^{-7} , respectively. The RME risk estimate is greater than, while the CT risk estimate is less than, the acceptable risk level of 1 x 10°s for cumulative carcinogenic risk, while the CT risk estimate is less than the DEQ acceptable risk level. The primary exposure pathways (RME evaluation) are soil ingestion (risk = 2 x 105) and dermal contact with soil (risk = 1×10^{5}). COPCs that exceed the DEQ acceptable risk level of 1×10^6 for individual carcinogens are benzo(a)pyrene (risk = 9×10^6) and arsenic (risk = 2×10^{-5}). The cumulative RME and CT HIs for the residential receptor are estimated to be 0.4 and 0.01, respectively. Both of which are less than the DEQ acceptable HI of 1.0. Hart Crowser Page 22 15191-01 January 18, 2002

Commercial Worker. The cumulative RME and CT excess lifetime cancer risks for the commercial worker are estimated to be 2×10^6 and 3×10^7 , respectively. The RME and CT risk estimates are both less than the DEQ acceptable level of 1 $\times 10^{-5}$ for cumulative carcinogenic risk. Arsenic (risk = 2×10^{-6}) is the only individual COPC that exceeds the DEQ acceptable risk-level of 1 x 106 for individual carcinogens. The cumulative RME and CT HIs for the residential receptor are estimated to be 0.01 and 0.006, respectively. The RME and CT HI are less than the DEQ acceptable HI of 1.0. Excavation Worker. The cumulative RME and CT excess lifetime cancer risks for the excavation worker are estimated to be 1×10^7 and 4×10^9 , respectively. The RME and CT risk estimates are less than the DEQ acceptable risk level for multiple carcinogens. The cumulative RME and CT HIs for the residential receptor are estimated to be 3×10^3 and 4×10^4 , respectively. Both of which are well below the acceptable HI-of-1.0. 3.3.2.3 Area C Risk and Hazard Estimates The exposure pathways that were quantitatively evaluated at Area C are soil ingestion, dermal contact with soil, and inhalation of fugitive dust. No VOCs were detected in Area C soil or groundwater. Arsenic is the only COPC for 'Area C. The cumulative RME and CT excess-lifetime cancer risks for all receptors (resident, commercial worker, and excavation worker) are less than the acceptable risk level of 1 x 10⁵ for cumulative carcinogenic risk, with the exception of the RME residential scenario (risk = 2 x 10⁻⁵). Arsenic has individual cancer risk estimates of 2 x 10⁻⁵ and 2 x 10⁻⁶ for the RME residential and commercial worker scenarios, respectively. All individual and cumulative hazard estimates are less than the DEQ acceptable HI of 1.0. 3.3.2.4 Lead Risk Evaluation Lead-was only identified as a COPC in Area A. Therefore, the discussion in this section only concerns Area A. Resident. The RME and CT lead EPCs in surface soil (0 to 3 feet bgs) are 6,190 mg/kg-and-540-mg/kg,-respectively.-Both-of-these-concentrations-exceed-the EPA Region 9 residential soil PRG of 400 mg/kg, indicating that there is a

A. The RME and CT lead EPCs in surface soil are driven by the maximum detected lead concentration of 6,190 mg/kg, which was detected in sample B-

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68. If sample B-68 was removed from the Area A surface soil data set, the RME and CT EPCs for the remaining data would be 192 mg/kg (based on the maximum detected concentration) and 30 mg/kg, respectively. Commercial Worker. The RME lead EPC in surface soil at Area A (6,190 mg/kg). exceeds, while the CT EPC is less than, the EPA Region 9 industrial soil PRG of 750 mg/kg. As discussed above, if the maximum detected concentrations were removed from the data set, the RME and CT lead EPCs would be acceptable for the commercial worker. Excavation Worker. The RME and CT lead EPCs in total soil (0 to 15 feet bgs) are 5,000 mg/kg and 430 mg/kg, respectively. The RME lead EPC exceeds, while the CT EPC is less than, the EPA Region 9 industrial soil PRG of 750 mg/kg. The RME and CT lead EPCs in total soil are also driven by the maximum detected lead concentration of 6,190 mg/kg that was detected in sample B-68. If sample B-68 were removed from the Area A total soil data set, the remaining maximum detected concentration would be 807 mg/kg, which just slightly exceeds the industrial soil lead PRG. 3.4 Uncertainty Analysis It is important to fully specify the assumptions and uncertainties inherent in the risk assessment to place the risk estimates in proper perspective. For this risk assessment, the general sources of uncertainty that are addressed include: Data collection-and evaluation: Exposure assessment; Toxicity assessment: and Risk characterization. 3.4.1 Data Collection and Evaluation The identification of the types and numbers of environmental samples, sampling procedures, and sample analysis each contain components that contribute to uncertainties in this risk assessment. For example, it is generally not practical to sample all locations and media at a site. Decisions were made to select a subset of the potential sampling locations and media-based upon the anticipated presence of the chemical. These decisions were made with the use of historical and background information of the site and the potential contaminants' chemical and physical properties. Exposure doses for the site that are based on non-random,-or-hot-spot,-samples-may be overestimated.

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3.4.2 Exposure Assessment The exposure estimation methods are subject to varying degrees of uncertainty: The degree of uncertainty generally depends on the amount of site-specific data available. The following sources of uncertainty have been identified. Exposure Scenario Identification. This HHRA assumes that receptors are limited to residents, commercial workers, and excavation workers. If this assumption is incorrect, future risks and hazards could be under- or overestimated. Exposure Parameters and Assumptions. The standard and site-specific exposure assumptions may or may not be representative of the actual exposure conditions and could under- or overestimate future risks and hazards. Calculation of Exposure Point Concentrations. The 90 percent UCL on the arithmetic mean, or the maximum detected concentration, whichever is lower, was used as the exposure point concentration (EPC) in this HHRA. Prior to the calculation of the 90 percent UCL, each data set was evaluated to determine whether the data were distributed normally or lognormally. As discussed previously, if a data set was found to be neither normal nor lognormal, the data set was evaluated as a lognormal data set. A lognormal distribution is common among environmental data sets. The maximum detected COPC concentrations, especially at Area A, has a significant effect on the EPCs used in this HHRA. Area A. Carcinogenic PAHs, arsenic and lead were identified as compounds of concern (COCs) in surface soil, while only lead was identified as a subsurface COC. The surface soil exposure concentrations for cPAHs, arsenic, and lead are driven by the maximum detected concentration of each COPC, which was detected in soil sample B-68 (B-94 for dibenz(a,h)anthracene). The second highest cPAH detections were found in soil sample B-94 (detected between 1 and 2 mg/kg). The remaining cPAH detections are less than 0.5 mg/kg, which are consistent with ambient levels of cPAHs in urban areas. Arsenic was detected at a concentration of 12.9 mg/kg in sample B-68 and at a concentration of 7.53 mg/kg in sample B-97. All other arsenic detections were less than the Terminal 1 arsenic background level of 5.3 mg/kg (Hahn and Associates, 2001a). Additionally, if the arsenic concentration of 12.9 mg/kg were removed from the Area A surface soil data set, the resulting arithmetic mean concentration of the remaining data would-be-2-2-mg/kg. As discussed previously, the maximum detected lead concentration in surface soil-at-Area A-was 6,190 mg/kg. The second highest lead detection was 192 mg/kg, which is below the residential and industrial (or commercial) Hart Crowser Page 25 15191-01 January 18, 2002

soil screening levels. Lead is the only COPC identified in the HHRA as a subsurface compound of concern (i.e., detected at levels above DEQ acceptable levels). Area B. Benzo(a)pyrene and arsenic were identified as COCs in surface soil. No subsurface COCs were identified at Area B. The maximum detected concentration of benzo(a)pyrene and the other three cPAHs evaluated are less than 0.2 mg/kg. These levels are consistent with ambient levels of cPAHs in urban areas. Arsenic was detected at a maximum concentration of 3.1 mg/kg in surface soil, which is less than the Terminal 1 arsenic background level of 5.3 mg/kg (Hahn and Associates, 2001a). Area C. Arsenic was identified as a COC in surface soil. No subsurface COCs were identified at AOC C. Arsenic was detected at a maximum concentration of 2.9 mg/kg in surface soil, which is less than the Terminal 1 arsenic background level of 5.3 mg/kg (HAI, 2001a). Assumption of Steady-State Conditions. The inherent assumption is that future COPC concentrations are the same as current concentrations. In general, this assumption overestimates COPC concentrations and resulting exposure intakes. Chemical Characterization. The sampling strategy used in collecting the soil samples that were used in this HHRA was purposive rather than random. Because the potential current and future receptors are assumed to visit the entire-site, not just the areas-that are contaminated, the exposure point concentrations used likely overestimate potential risks and hazards. Modeling Procedures. DEQ's Risk-based Decision Making guidance was used to estimate the volatilization from groundwater to indoor and outdoor air. The assumptions used in these models introduce uncertainty to the degree that they do not reflect actual conditions. There is significant uncertainty associated with the volatilization model used to estimate indoor and outdoor air concentrations based on soil and groundwater concentrations. Areas of uncertainty include, but are not limited to: ■ COPC Concentration: The model assumes the COPC concentrations are homogeneous over the entire area being evaluated. Since some COPC concentrations are based on the maximum detected concentration, this is a conservative assumption that is likely to significantly overestimate the amount of contamination present. Building Parameters: The model uses various building parameters as a basis for the indoor air concentrations such as building volume to area ratio Hart Crowser Page 26 15191-01- January-18, 2002

(essentially the height of the building), building air exchange rate (the amount of times the air in the building is replace per second), the foundation crack thickness, and the foundation crack fraction (that is, the fraction of the building floor that contains cracks). Many of these assumptions have a linear effect on the model output (that is, if the air exchange rate is doubled, the indoor-air concentration would drop in half). The model also assumes thereis no vapor barrier under the foundation and that the building is not under positive pressure. Default building parameters were used in this HHRA. **COPC-Specific Parameters:** The model uses various chemical parameters such as diffusion coefficients, Log K_∞ or Log K_d. Henry's Law Constant, vapor pressure, and solubility. These values can vary considerable in the literature. Default chemical parameters included in the RBDM model were used. These COPC-specific parameters can have a significant effect on the model results and, therefore, the degree that the parameters used represent actual conditions at the site may lead to an overestimation or underestimation of actual air concentrations. 3.4.3 Toxicity Assessment Whether verified by consensus among EPA scientists or not, uncertainty is present in the derivation of toxicity factors, and several assumptions are necessary. The factors used in the derivation of toxicity factors that add uncertainty to the results are presented below. Extrapolation from Animal Studies. Extrapolating human health risks from animal studies is complicated by physiological and pharmacokinetic differences. Similar toxic effects are not always observed in all species or at similar relative concentrations (when corrected for body weight). These extrapolations may overestimate or underestimate the actual chemical toxicity to humans. High-Dose to Low-Dose Extrapolations. Toxicity values are generally based on laboratory studies using high chemical exposures. Dose-response trends observed at high doses are generally assumed to be linear at low doses. Because dose-response relationships at low doses are largely-unknown, assuming a linear relationship may overestimate or underestimate chemical toxicity at concentrations in the extrapolated range. Population Variability. Laboratory animal studies generally use animal strains that are genetically similar, yet the human population is genetically diverse. Because methods for estimating toxicity in more susceptible individuals, such as children, are largely-undeveloped, such estimates may overestimate or underestimate chemical toxicity. Hart Crowser Page 27 15191-01 January 18, 2002

Available Studies. Not all toxicity values are based on the same amount or quality of research. As new studies are performed and reviewed, toxicity values can change. The less information available on a chemical, the greater the possibility that chemical toxicity will be overestimated or underestimated. The uncertainties discussed above are addressed when developing RfDs by dividing the no observable adverse effect level (NOAEL) from animal studies by uncertainty factors of up to 10,000. Uncertainty associated with determining chemical carcinogenicity is reflected in the weight-of-evidence classification groups assigned to carcinogens. In addition, uncertainties are introduced because SFs are derived from the lowdose end of the dose-response curves, and the experimental studies are usually conducted at the high-dose end of the curve. The selected 95 percent UCL of the slope of the dose-response curve is considered an upper-bound toxicity value. Therefore, it is unlikely that the SFs will underestimate risk. Actual cancer risk may range from a low of zero to the upper limit defined by the model. Uncertainty is also associated with using oral toxicity factors to evaluate dermal exposures. The use of oral toxicity factors as surrogates is necessary because there are no dermal toxicity factors approved by EPA. Most of the uncertainty exists because it is not known whether the compounds in question exhibit the same toxicity via dermal contact as they do via the oral pathway. Default oral absorption factors were used to adjust the oral toxicity factors so that the absorbed doses calculated for the dermal pathway could be evaluated. The use of the oral absorption factors may bias the risk and hazard estimates high or low. The use of surrogate toxicity factors for chemicals lacking toxicity factors may under- or overestimates the potential risks or hazards. 3.4.4 Risk Characterization This HHRA used EPA's standard algorithms to calculate chemical intakes and associated health risks and hazards. There are certain assumptions inherent in the use of these equations that add uncertainty. For example, calculations of carcinogenic risks and non-carcinogenic HIs assume the additivity of toxic effects. This assumption adds uncertainty to the assessment and may result in an overestimation or underestimation of the potential risks, depending on whether synergistic or antagonistic conditions apply. Exposure pathway risks are combined assuming that a single receptor may be exposed to contamination through a selected number of pathways concurrently. This is a conservative estimate that may overestimate risks and hazards. Additionally, the standard algorithms used do not consider certain factors, such as absorption or matrix Hart Crowser Page 28

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	· · · · · · · · · · · · · · · · · · ·
	effects. In cases where these processes are important, the risk estimates may
	overestimate or underestimate the potential human risks at this site.
	over standard over an action of the potential region at the sales
4.0 LE\	/EL 1 SCOPING ECOLOGICAL RISK ASSESSMENT
	The purpose of the Level 1 Scoping ERA is to provide a qualitative determination
	of whether there is any reason to believe that ecological receptors and/or
	exposure pathways are present or potentially present at or in the locality of the
	facility. The outline for the Level 1 Deliverable (Ecological Risk Assessment
	Guidance; Attachment 3, DEQ, 1998) was generally followed for presenting the
	results of the Level 1 evaluation in this section. The existing data summary and
· · · · · · · · · · · · · · · · · · ·	the results of the land and water use survey are presented in earlier sections of
,	this report. Appendix C presents photographs taken at the T1S Site during the
	site visit and Appendix D presents DEQ's Ecological Scoping Checklist.
	In addition to the Level 1 Scoping ERA, a Modified Level 2 Screening ERA was
	conducted on the groundwater data available for this site. The Modified Level 2
	Screening ERA was conducted to determine whether constituents were present
	in groundwater at levels of concern for aquatic ecological receptors.
	The site and expressed in a proportion are all a made becausing district and are
	The site and surrounding properties are all zoned heavy industrial and are
	being used for these purposes. The Willamette River borders the T1S Site on
	the east. The T1S Site has been historically used for commercial use and the
	entire site has been developed. The site does not provide high quality habitat
<u> </u>	to the local ecological community. There are no designated wetlands on the locality of the facility. There are also no identified sensitive environments in
•	the locality of the facility.
	the locality of the facility.
4.2 Thr	eatened and Endangered Species
-	The Oregon Natural Heritage Program (ONHP), which monitors
	threatened and endangered (T&E) plants and wildlife, conducted a data
	search of T&E species within a two-mile radius of the site. A letter from
	the ONHP is included in Appendix E. The ONHP identified the historical
	presence of the following species:
·	Federal Species Listed as Threatened
	Oncorhynchus mykiss (Steelhead-[I:ower-Columbia-River and Upper
	Willamette River]).
	(Vindividue Niverij).
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·- ·-	
	 Oncorhynchus tshawytscha (Chinook salmon [Lower Columbia River and
	Upper Willamette River]).
	·
"	Candidate for Federal Listing as Threatened
	■ Oncorhynchus kisutch (Coho salmon [Lower-Columbia River]).
_	
	Federal Species Listed as Proposed Threatened
B	 Oncorhynchus clarki (Coastal cutthroat trout [Columbia River/SW
	Washington).
	Federal Species of Concern
	■ Corynorhinus townsendii townsendii (Pacific Western Big-Eared Bat) — Last
	observed-in-1928.
· ·	
	 Antrozous pallidus pacificus (Pacific Pallid Bat) – Last observed in 1927.
	■ Clemmys marmorata marmorata (Northwestern Pond Turtle).
•	
.	 Aster Curtus (White-Topped Aster) - This population is assumed extinct.
	State Species Listed as Endangered
	State species Eisted as Endangered
	Falco peregrinus anatum (American Peregrine Falcon) - Nesting observed
· · · · · · · · · · · · · · · · · · ·	in 1997.
	State Species Listed as Sensitive-Critical
	Chrysemys picta (Painted Turtle).
•	
,	State Species Listed as Critical
	- Children - 1 - 17-110 - 1 - 1 - 1000
_	■ Cimicifuga elata (Tall Bugbane) – Last observed in 1993.
	T&E species were not observed in the upland portions of this site on our
, ,	ecological scoping visit.
-	
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	8, 2002

4.3 Site Visit Summary

This section describes the results of Hart-Growser's October 2, 2001, visit to the site to assess whether ecological receptors and/or exposure pathways are present or potentially present at or in the locality of the site. The discussion of ecological features present at the facility is based on our on-site observations.

Photographs taken during the site visit are provided in Appendix C.

4.3.1 Observed Impacts

Impacts to the site and surrounding properties attributable to contaminated environmental media were not observed. The entire site has been developed for commercial and industrial use; as such, native vegetation has been replaced with buildings and pavement.

4.3.2 Ecological Features

Ecological features were assessed by evaluating the habitat within the locality of the facility. Appendix D presents-the-checklists-used-in-this evaluation.

Upland. The T1S Site consists of approximately 21-acres of flat terrain with limited on-site vegetation. The site is 99 percent ruderal and 1 percent vegetated (Figure 2 and Photos in Appendix C). Upland vegetation is limited to a small bank area that slopes immediately bordering the Willamette River. The types of vegetation observed in this bank area were limited to invasive, weedy species such as Blackberry (*Bubus sp.*). Additionally, limited vegetation was observed in cracks in the pavement throughout the site. In general, the entire upland site has been paved or developed and provides very poor habitat quality and extremely limited potential for exposure to terrestrial ecological receptors.

4.3.3 Ecologically Important Species and Habitats

Ecologically important terrestrial species, including threatened or endangered animals were not observed on or adjacent to the site.

4.4 Exposure Pathways

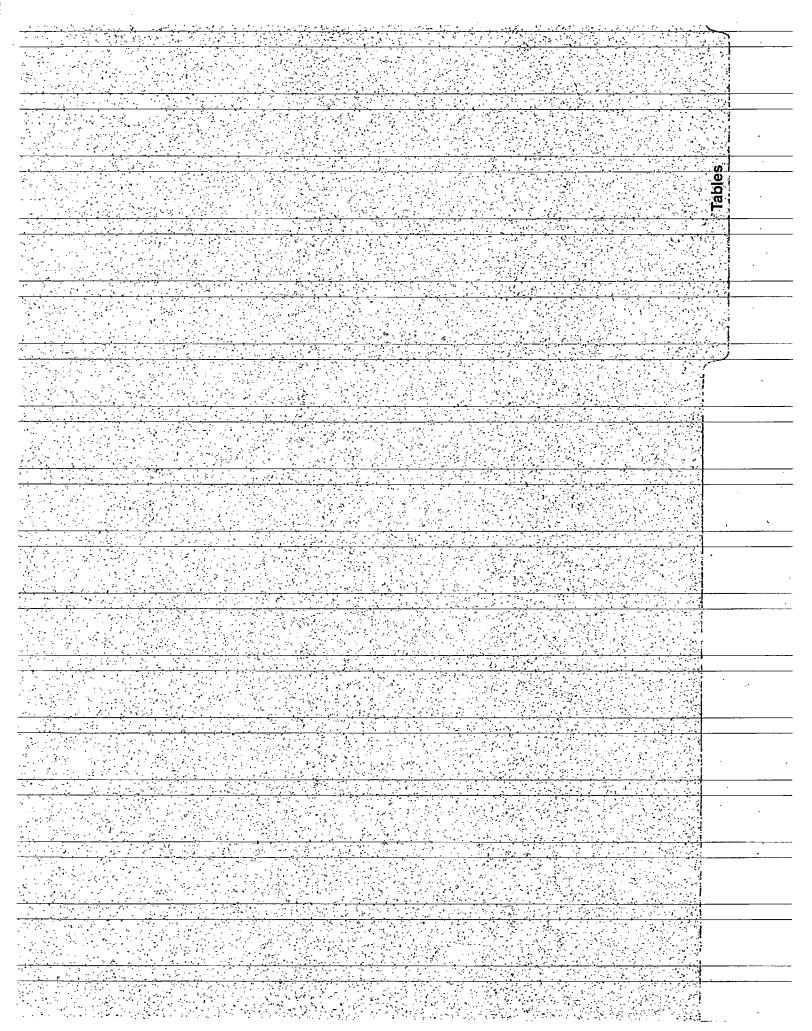
A general evaluation of potential receptor-pathway interactions is provided in the checklists presented in Appendix D and is presented in the Ecological CSM—on-Figure 4.—As-summarized-on-the-checklists provided-in-Appendix-D, GOPGs-are currently present in soils within the locality of the facility. However, there are no current exposure pathways present for these contaminants to reach ecological receptors within the locality of the facility. The majority of the site is currently paved or is covered by buildings. The absence of upland habitat

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results in no complete exposure pathways to terrestrial species. The fact that the majority of the site is paved or developed limits the potential for overland transport (via surface erosion of soil) to cause migration of any contamination present in surface soil to the Willamette River. The RI found no evidence of free phase product to be present in soil, and no migration or direct release of product from the T1S Site to the adjacent Willamette River is expected at this site. The Preliminary Assessment (PA) completed for this site by the Port (Port of Portland, 2000) evaluated the potential for storm water discharges to transport contaminants from the T1S Site to the Willamette River. The PA concluded that "there is also a low potential for upland activities to have resulted in releases to the Willamette River via stormwater discharges". Groundwater data collected from push-probe explorations as part of the RI suggested the potential for site related contaminants to be present in site groundwater. A groundwater monitoring program was initiated by Hahn and Associates and seven monitoring wells were installed at the T1S Site (Hahn and Associates, 2001b). Because the movement of shallow groundwater at the site has been found to be in the direction of the Willamette River, a modified Level 2 Screening ERA was conducted on the available groundwater monitoring data to determine whether contaminants are present in groundwater at concentrations at levels of potential concern to ecological receptors. The procedures for conducting a Modified Level 2 Screening ERA were presented in the Risk Assessment Work Plan for this site (Hart Crowser, 2001) and further discussed with DEQ in Port of Portland's Response to Review Comments letter (Port of Portland, November 12, 2001). The available groundwater monitoring data were screened against appropriate DEQ Ecological Screening Benchmark Values (SBVs) to determine whether the detected concentrations of contaminants exceeded the risk based screening levels. The Modified Level 2 Screening of groundwater data is presented in Table 12. The groundwater monitoring well data from each well-were-screened against the Freshwater Aquatic SBVs. No PAHs or VOCs were detected in groundwater at concentrations exceeding their corresponding SBVs. There were two metals (lead and mercury in MW-3 and lead in MW-7) for which the total metal concentration-exceeded the corresponding SBV. However, the analysis of the dissolved fraction of lead and mercury from monitoring wells MW-3 and MW-7 indicates these metals were not detected in the samples and there are no detected concentrations of analytes that exceed the SBVs. As dissolved metals represent the bioavailable fraction of metals in aqueous media, it is concluded Hart Crowser Page 32 15191-01 January 18, 2002

	that there are no constituents in groundwater at levels of concern to aquatic
	ecological receptors.
4.5 Concl	lusions and Recommendations
	In October 2001, Hart Crowser completed a Level 1 Scoping ERA for possible
•	ecological receptors and pathways at the T1S Site. The site visit and historical
	research did not identify any ecologically important species or habitats present in
	the upland portion of this site. The site is almost entirely paved or covered by
	buildings. The absence of upland habitat results in no complete exposure
·	pathways to terrestrial species.
	A Modified Level 2 Screening ERA was conducted on the available groundwater
	data collected at this site. There were no detected concentrations of organic
	constituents in the seven groundwater monitoring wells that exceeded their
	corresponding Ecological-SBVs. There were two metals (copper and lead)
<u></u>	detected in groundwater that exceeded SBVs based on the analysis of unfiltered,
	total metals but when the same samples were analyzed for dissolved metals,
	copper and lead were not detected. The dissolved fraction of metals represents
	the bioavailable fraction in aqueous environmental media. Therefore, it is
	concluded that there is no potential for adverse ecological impacts to aquatic
	ecological receptors from the discharge of groundwater to the Willamette River.
	No-further ERA activities are warranted at this site.
	No further EXA activities are warranted at this site.
•	
5.0 LIMIT	ATIONS
	Hart Crowser performed this work in accordance with generally accepted
	professional practices related to the nature of the work accomplished, in the
·-···	same or similar localities, at the time the services were performed. This report is
•	for the specific application to the referenced project and for the exclusive use of
	the Port. No other warranty, express or implied, is made.
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	· · · · · · · · · · · · · · · · · · ·
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Tabl	e 1	- Area A (O	C Id	entifi	cat	on Table	
Mari	ne	Terminal	t S	outh	Risk	As	sessment	l
Port	Jan	d, Oregon	1					l

Total Petroleum Hydrocarbons Diesel Dil Somivolatiles	PRG NA NA 3.7E+03	Cij 1.2E+03 1.8E+03	Concentra Rij	COPC? Yes¹	q/kg) Rij/Rj	COPC?	PRG	Ground Cij	vater (Conc Rij	COPC?	RIVRI	COPC?	SRIJ	Multiple Media COPC?
Diesel Dil Somivolstiles	NA NA 3.7E+03	1.2E+03	-		RijiRj	COPCY	PRG	Cij	KI N	COPC7	RIVRI	COPCY	SKI	COPC?
Diesel Dil Somivolstiles	NA 3.7E+03			Yes¹										
Oil Somiyolatiles	NA 3.7E+03			Yes'_		·	ļ., ļ.,						<u> </u>	
Somivolatiles	3.7E+03	1,86+03				Yes'	NA NA	4.2E+02	-	Yes¹	<u>. - </u>	Yes'	กล	No
		1		Yes'		Yes'		ND		-	-	-	na	lNo_
Acenaphthene		6.6E+00	1,8E-03	No	8.95-06	No	3.7E+02	7.2E-01	1.9E-03	No	7.6E-04	No	3.7E-03	Nο
Acchaphthylene	3.7E+03	5.6E-01	1.5E-04	No	7.65-07	No]	ND		<u> </u>	_		1.5E-04	No
Anthracens	2.2E+04	1.1E+01	5.2E-04	No	2.6E-06	No	1-	79					5.2E-04	No
Berzo(a)anthracene	6.2E-01	9.4E+00	1.5E+01	Yes	7.6E-02	Yes	(I-	(4)		T -		- 1	1 5E+01	Yes
Benzo(a)pyrene	6.2E-02	7.1E+00	1.1E+02	Yas	5.7E-01	Yos	III-	70	-		- 1		1.1E+02	Yes
Benzo(b)(kuoranthene	6.2E-01	4.2E+00	6.8E+00	Yes	3.4E-02	Yes	I -	ND	-	 -		1	8 8E+00	Yes
Benzp(g.h,i)perylene	2.3E+03	3.8E+00	1.6E-03	No	8.2E-06	No	-	ND		-		- 1	1,6E-03	No
Benzo(k)fluorantnene	8.2E+00	5.5E+00	8.9E-01	No	4.5E-03	No	-	ND		-	- 1	-	8.9E-01	No
Chrysene	6.2E+01	9.6E+00	1.5E-01	No	7.8E-04	No	<u> </u>	ND		7-	-		1.5E-01	No
Diberz(a,h)anthracene	6.2E-02	3.5E-01	5.6E+00	Yes	2.8E-02	No		70	-	-			5 6E+00	Yes
luoranthene	2.3E+03	2.0E+01	8.5E-03	No	4.3E-05	No	-	NE		_			8.5E-03	No
enenoul	2.6E+03	5,7E+00	2.2E-03	No	1,1E-05	No	I -	ND		—			2.2E-03	No
ndeno(1.2.3-cd)pyrene	8.2E-01	8.4E+00	5.5E+00	Yes	2.7E-02	No		ND		T-	-		5 5E+00	Yes
Vaphthalene	5.6E+01	7.9E+00	1.4E-01	No	7.1E-04	No	6.2E+00	2.9E-01	4.7E-02	l No	1,8E-02	No I	1.9E-01	No
Phenanthrene	2.2E+04	3.5E+01	1.6E-03	No	8.05-08	No	1.9E+03	1.3E+00	6.9E-04	No	2.7E-04	No I	2.3E-03	No
Pyrene	2.35+03	2.8E+01	1.2E-02	No	6.0⊒-05	No	1 - T-	ND		1 -	1-1		1.25-02	No
Metals							1							
Antinjonyi I	3.1E+01	2.0E+01	6.3E-01	No	3.2E-03	No		ND I	_	-	1-1-		6.3E-01	No
Arsenic	3.9E-01	1.3E+01	3.3E+01	Yes	1.7E-01	Yes	-	NE	1	-	<u> </u>		3 3E+01	Yes
Cadmium I	3.7E+01	7.4E+00	2.0E-01	No	1.0E-03	No		- ND		T-	<u> </u>	-	2.0E-01	No
Chromium	2.1E+02	4.3E+01	2.1E-01	No	1.0E-03	No	-	ND		-	1_	-	2.1E-01	No
Copper	2.9E+03	2.9E+02	9.9E-02	No	5.0E-04	No	-	NE			-		9.9E-02	No
_ead	4.0E+02	6.2E+03	1.5E+01	Yes	7.8E-02	Yes		NE	-	-	<u> -</u>	-	1!5E+01	Yes
Mercury	2.3E+01	9.7E+00	4.1E-01	No	2.1E-03	No	_	ND	-		-		4.1E-01	No
Vickel	1.6E+03	6.4E+01	4.0E-02	· No	2.0⊟-04	No	_	ND		-	-	_	4.0E-02	No
Thatium	5.2E+00	3.5E+00	8.7E-01	No	3.4E-03	No	-	ND	-	-		-	6.7E-01	No
Zinc	2.3E+04	3.0E+02	1.3E-02	No	6.66-05	No	-	ND				-	1.3E-02	No
Volatiles	1			 	1	<u> </u>		1	1					
Acetone	1.6E+03	2.2E-01	1.4E-04	No	7.16-07	No	1 -	ND		-			1.4E-04	No
2-Butanone	7.3E+03	3.6E-02	4.7E-08	No	2.4E-08	No	-	ND			-	-	4.7E-08	No
Tetrabhloroethene .	11 -	nd	-	No	-	No	1.1E+00	2.8E+00	2.5E+00	Yes	9.8E-01	Yes	2,5E+00	Yes
RI			2.0E+02		•	· · · · · · · · · · · · · · · · · · ·			2.65+00					
NI	1-1-		3.0E+01	T					5.0⊆+00			·		
1/Nij			3.3E-02	 		·	1 		2.0E-01		. 11	Ì	į	1 1

F:\DATA\Jobs\Port of Portland\15181-01 T-1 Risk Assessment\Tables\Tables 1-3, COPC

Notes:

(1) COPC identified based on the presence of TPH in site soils and groundwater. No PRG is available for screening.

Variables:

Variables:

PRG = EPA Region 9 PRG (residential for soil, tap water for groundwater).

Cij ≈ Maximum detected concentration of compound I in medium J.

Rij ≈ Risk ratio for compound I in medium J. (Cil/PRG); compound is a COPC if Rij is greater than 1.

Rij ≈ Sum of risk ratios for medium J.

Nij ≈ Number of compounds I detected in medium J.

Rij/Rij ≃ Compound is a COPC if this ratio is greater than 1/Nij.

SRij ≈ Summery risk ratio for compound I in at media (total Rij across all media); compound is a COPC if Srij is greater than 1.

Acronyma: NA = Not Available. ND = Not Detected.

NE = Not Evaluated (only volate compounds evaluated).

- = Not Applicable.

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Tabl	e 2	- Area	BC	OP:	C lo	lent	fic	ation Tal	le	I
Mari	пө	Termir	nal 1	\$o	uth	Ris	k A	ssessme	nt	
Port	lan	d, Ores	gon	1						Ì

		Soll	Concentre	tions in m	g/kg)			Ground	vater (Cond	entrations	in µg/L)		l T	Medium
	PRG	CIJ .	Rij	COPC?	Rij/Rj	COPC?	PRG	Cij	Rij	COPC?	Rij/Rj	COPC?	SRIJ	COPC?
Total Petroleum Hydrocarbons														
Diesel I	NA.	1.2E+03		Yes'	-	Yes	NA	4.2E+02	-	Yes	_	Yes	na	No
OII I	NA.	1.8E+03		Yes ¹	—	Yes		_	-	-			ла	No
Semivolatiles		1					1		 - - - 			l	 	
Acenaphthene	3.7E+03	1.1E+02	2.9E-02	No	4.8E-04	No	3.7E+02	7.2E-01	1.9E-03	No	-	-	3.1E-02	No
Acenaphthylene	13.7E+03	2.6E-01	7.1E-05	No	1.2E-06	No	1 1	ND	<u> </u>	1	-1		7.1E-05	No
Anthracene	2.2E+04	6.8E+01	3.1E-03	No	5.2E-05	No		ND	-			_	3.1E-03	No
Benzo(a)anthracene	6.2E-01	1.5E+00	2.4E+00	Yes	4.1E-02	Yes	1	ND	-		-	-	2.4E+00	Yes
Benzo(a)pyrene	6.2E-02	2.4E+00	3.8E+01	Yes	6.3E-01	Yes		ND	-				3.8E+01	Yes
Benzo(b)fluoranthene	6.2E-01	1.5E+00	2.5E+00	Yes	4.2E-02	Yes	1 4	ND	-	1			2.5E+00	Yes
Benzo(g,h,i)perylana	2.3E+03	2.5E+00	1.1E-03	No	1.8E-05	No	1 4 1	ND	_	1	-1		1,1E-03	No
Benzo(k)fluoranthene	6.2E+00	1.1E+00	1.8E-01	No	2.9E-03	No	1	ND	-	11	_	-	1,8E-01	No
Chrysene	6.2E+01	1.9E+00	3.0E-02	No	5.1E-04	No	1 4	ND	-				3,0E-02	No
Diberiz(a,h)anthracene	6.2E-02	2.5E-01	4.1E+00	Yes	6.8E-02	Yes	4	ND				_	4.1E+00	Yes
luoranthene	2.3E+03	2.9E+02	1.2E-01	No	2.1E-03	No	1 + 1	NE	_			-	1,2E-01	No
luorene	2.6E+03	1.7E+02	6.7E-02	No I	1.1E-03	No	1	ND	-				6.7E-02	No
ndeno(1,2,3-cd)pyrene	6.2E-01	1.6E+00	2.5E+00	Yes	4.2E-02	Yes	1 4	ND	_	-1		_	2.5E+00	Yes
Naphthalene I ;	5.6E+01	4.0E+00	7.1E-02	No	1.2E-03	No	8.2E+00	2.9E-01	4.7E-02	No		-	1,2E-01	No
Phenanthrene	2.2E+04	7.0E+02	3.2E-02	No	6.3E-04	No ·	1.8E+03	1.3E+00	6.9E-04	No	-	_	3.3E-02	No
Pyranie	2.3E+03	1.4E+02	6.2E-02	No	1.0E-03	No		ND	_	1-	-	_	6.2E-02	No
Metals														
Antimony	3.1E+01	7.0E-01	2.3E-02	No	3.8E-04	No		ND	-	-		-	2.3E-02	No
Arsenic	3.9E-01	3.6€+00	9.2E+00	Yes	1.5E-01	Yes		NE	-	-	-		9.2E+00	Yes
3eryllium	1.5E+02	2.3E-01	1.5E-03	No	2.6E-05	No	4	ND I	-	-	-	_	1.5E-03	No
Cadmium	3.7E+01	1.3E+00	3.6E-02	No	6.1E-04	No	1	ND	_				3.6E-02	No
Chromium	2.1E+02	1.6E+01	7.8E-02	No	1.3E-03	No	4	ND		-			7.8E-02	No
Copper	2.9E+03	2.7E+01	9.3E-03	No	1,8E-04	No	1 4 1	NE		~	9-7	_	9.3E-03	No
bse	4.0E+02	1.2E+02	3.0E-01	No	5.1E-03	No		NE	-] 	3.DE-01	No
Mercury	2.3E+01	1.5E-01	6.2E-03	No	1.0E-04	No	-	ND	-		-		6.2E-03	No
vickel	1.6E+03	2.4E+01	1.5E-02	No	2.5E-04	No		ND	_	1		_	1.5E-02	No
[hallium]	5.2E+00	6.1E-02	1.2E-02	No	2.0E-04	No	_ +	ND			-		1.2E-02	No
Zine	12.3E+04	1.4E+02	6.1E-03	No	1.0€-04	No		ND		-		-	6.1E-03	No
RJ			6.0E+01						5.0E-02					
Nij	·	11	2.9E+01						4,0E+00					
1/NI)		1 1 1 1 1	3 45-02	1					2.5F-01	1 1 1	1	1	1 1	1 1

P\DATAUcbs\Fort cf Portland\15191-01 T-1 Risk AssessmennTables\Tables 1-3, COPC

Notes:

(1) COPC identified based on the presence of TPH in site soils and groundwater. No PRG is available for screening.

Variables:

PRG = EPA Region 9 PRG (residential for soil, tap water for groundwater).

Cij = Maximum detected concentration of compound I in medium i.

Rij = Risk ratio for compound I in medium j (Cij/PRG); compound is a COPC if Rij is greater than 1.

R) = Sum of risk ratios for medium j.

Nij = Number of compounds i detected in medium j.

Rij = Compound is a COPC if this ratio is greater than 1/Nij.

SRij = Summary risk ratio for compound in all media (total Rij across all media); compound is a COPC if Srij is greater than 1.

Adronyms: NA = Not Available.

ND = Not Detected.

NE = Not Evaluated (only voletile compounds evaluated).

- Not Applicable.

Table 3 - Area C COPC Identification Table Marine Terminal 1 South Risk Assessment Portland, Oregon

		Soil	(Concent	tions in m	gkg)			Ground	water (Con	centrations	ln μg/L)			Mediun
	PRG	Cij	RIJ	COPC?	Rij/Rj	COPC?	PRG	Cij	Ri)	COPC?	Rij/Rj	COPC?	SRIJ	COPCT
Metals										11				
Arsenic	3.9E-01	1.2E+01	3.0E+01	Yes	9.9E-01	Yes		-	-	11 - 1	-	- -	3.0E+01	Yes
Beryllium	1.5E+02	7.8E-01	5.2E-03	No	1.7E-04	No		-	-	T -	_		5.2E-03	No
Cadmium	3.7E+01	1.3E+00	3.5E-02	No	1.2E-03	No	-			11 - 1	<u>.</u>	 	3.5E-02	No
Chromium I	2.1E+02	2.6E+01	1.2E-01	No	4.0E-03	No	-			-	-	-	1.2E-01	No
Copper	2.9⊑+03	1.9E+01	6.6E-03	No	2.1E-04	No			-	_	-	11-	6.6E-03	_ No
Lead	4.0E+02	1.6E+01	3.9E-02	No	1.3E-03	No	_	-		-			3.9E-02	No
Nickel	1.6E+03	1.9E+01	1,2E-02	No	3.9E-04	No				-	_		1.2E-02	No
Zinc I	2.3E+04	7.7E+01	3.3E-03	No	1.1E-04	No			- 1		_		3.3E-03	No
Rj l			3.0E+01						0.0E+00					
NIJ			8.0E+00						0.0E+00	1		1 1	i i) '
1/NIj		i i	1.3E-01		11					7	1	11	1 1	•

F: DATA Jobs Port of Portand 15191-01 T-1 Risk Assessment Tables Tables 1-3, COPC

Acronyms:

- = Not Applicable.

Variables:

PRG = EPA Region 9 PRG (residential for soil, tap water for groundwater).

Cij = Maximum detected concentration of compound i in medium j.

RIJ = Risk ratio for compound i in medium j (dij/PRG); compound is a COPC if Rij is greater than 1.

Rj = Sum of risk ratios for medium j.

Nij = Number of compounds i detected in medium j.

Rij/Rij = Compound is a COPC if this ratio is greater than 1/Nij.

SRij = Summary risk ratio for compound i in all media (total Rij across all media); compound is a COPC if Srij is greater than 1.

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Table 4 - Exposure Point Concentrations: Soil and Groundwater Marine Terminal 1 South Risk Assessment Portland, Oregon

		<u> </u>					1-1	EP	C
Analyte	Detection Frequency	SQL Range (Min-Max)	Detect Range (Min-Max)	Sample ID of Maximum Detection	Distribution	90 % UCL	Arithmetic Mean	RME	СТ
AREA A: SURFACE SOIL	(0 to 3 feet by	gs)							
PAHs in mg/kg			1						
Benzo(a)anthracene	7/18	0.0134 - 1.0	0.0275 - 9.35	B-68	Weak Lognormal	2 0E+00	6.7E-01	2.0E+00	6.7E-01
Benzo(a)pyrene	7/18	0.0134 - 1.0	0.0292 - 7.05	B-68	Weak Lognormal	1,8E+00	5.5E-01	1.8E+00	5.5E-01
Benzo(b)fluoranthene	6/18	0.0134 - 1.0	0.0189 - 4.22	B-68	Weak Lognormal	1 4E+00	4.0E-01	1.4E+00	4.0E-01
Dibenz(a,h)anthracene	3/18	0.01 - 1.34	0.033 - 0.16	B-94	Weak Lognormal	1.8E-01	9.0E-02	1.6E-01	9.0E-02
Indeno(1,2,3-cd)pyrene	6/18	0.0134 - 1.0	0.0131 - 3.3B	B-68	Weak Lognormal	7.4E-01	2.8E-01	7.4E-01	2.8E-01
Metals in mg/kg									
Arsenic	7/10		1.49 - 12.9	B-68	Lognormal	8.4E+00	3.3E+00	8.4E+00	3.3E+00
Lead	7/12	1.0 - 10.0	9.21 - 6,190	B-68	Lognormal	1.7E+04	5.4E+02	6.2E+03	5.4E+02
TPH in mg/kg									
Dlesel Range	5/18	24.4 - 506	36.0 - 653	B-68	Assm. Lognormal	1.8E+02	9.7E+01	1.8E+02	9.7E+01
Oil-Range	6/18	50.0 - 61.7	72.1 - 1,300	B-94	Assm. Lognormal		2.1E+02	3.8E+02	2.1E+02
AREA A: TOTAL SOIL (0 t	o 15 feet bgs								
PAH's in mg/kg									
Benzo(a)anthracene	16/41	0.01 - 1.0	0.0203 - 9.35	B-68	Weak Lognormal	3.5E-01	3.7E-01	3.7E-01	3.7E-01
Benzo(a)pyrene	16/41	0.01 - 1.0	0.0157 - 7.05	B-68	Weak Lognormal	3.7E-01	3.3E-0.1	3.7E-01	3.3E-01
Benzo(b)fluoranthene	16/41	0.01 - 1.0	0.018 - 4.22	B-68	Weak Lognormal	3.4E-01	2.7E-01	3.4E-01	2.7E-01
Dipenz(a,h)anthracene	6/41	0.01 - 1.34 [0.015 - 0.35	B-38	Weak Lognormal	7.0E-02	6.0E-02	7.0E-02	6.0E-02
Indeno(1,2,3-cd)pyrene	14/41	0.01 - 1.0	0.0131 - 3.38	B-68	Weak Lognormai	2.0E-01	1.8E-01	2.0E-01	1.8E-01
Metals in mg/kg	! !	1		1 1 1		l 1	1		
Arsenic	12/15		1.35 - 12.9	B-68	Lognormal	6.0E+00	3.4E+00	6.0E+00	3.4E+00
Lead	11/18	10-10.0	2.73 - 6,190	B-68	Lognormal	5 0E+03	4.3E+02	5.0E+03	4.3E+02
TPH in mg/kg									
Diesel Range	11/53	24.4 - 506	25.5 - 1,170	B-102	Assm. Lognormal	8.7E+01	9.6E+01	9.6E+01	9.6E+01
Oil-Range	16/53	50.0 - 69.4	62.0 - 1.760	B-102	Assm. Lognormal	18E+02	2.0E+02	2.0E+02	2.0E+02
AREA A: GROUNDWATER		·		'					
Tetrachloroethene	1/2	1.0	2.76	MW-1	NA .	2.8E+00	2.8E+00	2.8E+00	12.8E+00
Please refer to notes at e	nd of table.				<u>' </u>	1		· · · · · · · · · · · · · · · · · · ·	

								EP	С
Analyte	Detection Frequency	SQL Range (Min-Max)	Detect Range (Min-Max)	Sample ID of Maximum Detection	t	90 % UCL	Anthmetic Mean	RME	СТ
AREA B: SURFACE SOIL	. (0 to 3 feet b	gs)							
PAHs in mg/kg									
Benzo(a)anthracene	4/7	0.0134 - 0.067	0.0448 - 0.149	B-87	Løgnormal	1.7E-01	6.0E-02	1.5E-01	6.0E-02
Benzo(a)pyrene	4/7	0.0134 - 0.067	0.0501 - 0.188	B-87 .	Lognormal	2.1E-01	7.0E-02	1.9E-01	7.0E-02
Benzo(b)fluoranthene	4/7	0.0134 - 0.067	0.0402 - 0.134	B-64/B-64a	Lognormal	1.8E-01	6.0E-02	1.3E-01	6.0E-02
Indeno(1,2,3-cd)pyrene	4/7	0.0134 - 0.067	0.0281 - 0.117	B-64/B-64a	Lognormal	1.4E-01	5.0E-02	1.2E-01	5.0E-02
Metals in mg/kg					·			'	1
Arsenic	4/4		2.6 - 3.1	B-31	NA I	3.1E+00	2.9E+00	3.1E+00	2.9E+0
TPH in mg/kg		1 1 1							
Oil-Range	4/9	50.0	1,170 - 6,030	B-5	Weak Lognormal	1.6E+05	1.2E+03	6.0E+03	1.2E+0
AREA B: TOTAL SOIL (0	to 15 feet bgs	3)							I
PAHs in mg/kg									ii
Benzo(a)anthracehe	13/19	0.0134 - 67	0.02 - 1.51	B-92	Lognormal	1.7E+00	4.0E-01	1.5E+00	4.0E-01
Benzo(a)pyrene	13/19	0.0134 - 67	0.0246 - 2.35	B-92	Lognormal	2.1E+00	4.7E-01	2.1E+00	4.7E-01
Benzo(b)fluoranthene	13/19	0.0134 - 67	0.0215 - 1.54	B-92	Lognormal	1.4E+00	3.7E-01	1.4E+00	3.7E-01
Diberz(a,h)anthracene	4/19	0.0134 - 67	0.0839 - 0.253	B-92	Lognormal	2.3E-01	1.1E-01	2.3E-01	1.1E-01
Indeno(1,2,3-cd)pyrene	13/19	0.0134 - 67	0.0193 - 1.56	B-92	Lognormal	9. 0E- 01	2.9E-01	9.0E-01	2.9E-01
Metals in mg/kg	j	1						1	
Arsenic	6/6		2.05 - 3.6	B-63/B-63a	Lognormal	3.3E+00	29E+00	3.3E+00	2.9E+00
TPH in mg/kg								1	
Diesel Range	5/30	20.8 - 2,500	29.0 - 3.440	B-63/B-63a	Assm. Lognormal	4.8E+02	27E+02	4.8E+02	2.7E+02
Oil-Range	16/30	50.0 - 52.1	62.7 - 20,700	B-65/B-65a	Assm. Lognormal	6!8E+03	1.7E+03	6.8E+03	1.7E+0
AREA C: SURFACE SOIL	(0 to 3 feet b					1.		1	
Metals in mg/kg						T			
Arsenic	1/1		2.9	8-32	NA	2.9E+00	2.9E+00	2.9E+00	2.9E+0
AREA C: TOTAL SOIL (0	to 15 feet bgs	3)				1			
Metals in mg/kg									
Arsenic	3/3		2.72 - 11.8	B-3	NA NA	1.2E+01	5.8E+00	1.2E+01	5.8E+0
		' - - '			DATAWobs\Port of Por	tland\15191-01		mentTablesT	able 4. EP

Notes:

Acronyms and Abbreviations:

EPC = Exposure point concentration.

PAHs = Polynuclear aromatic hydrocarbons.

RME = Reasonable maximum exposure.

TPH = Total petroleum hydrocarbons.

UCL = Upper confidence limit on the mean.

VOCs = Volatile organic compounds.

CT = Central Tendency.
SQL = Standard quantification limit.
NA = Not applicable.

Table 5 - Exposure Dose Equations and Exposure Factor Values: Soil Ingestion
Marine Terminal 1 South Risk Assessment
Portland, Oregon

$LADD^{a}(mg/kg-d) = \underline{C}_{so}$	R X IRS X CF X EF X ED BW X Atom
$ADD^{b}(mg/kg-d) = \underline{C}_{so}$	3 × IRS × CF × EF × ED BW × At _{ron}
EXPOSURE FACTOR (units)	RME ^a Value CT ^f Value
C _{soil} = Chemical concentration in soil (mg/kg)	UCL ₉₀ ^c Arithmetic Mean
CF=Conversion factor (kg/mg)	10-8 10-6
IRS = Incidental Soil Ingestion Rate (mg/d) Commercial Worker	100 ^d 50 ^d
Utility/Excavation Worker	480 ^d 100 ^d
Resident – Adult Resident – Child	100 ^d 50 ^d 400 ^d 100 ^d
EF = Exposure frequency (days/year)	
Commercial Worker	250 ^d 250 ^d
Utility/Excavation Worker	9 ^d 9 ^d
Resident – Adult/Child	350 ^d 40 ^d
ED = Exposure duration (year)	
Commercial Worker	25 ^d 6 ^d
Utility/Excavation Worker	1 ^d 0.5 ^d 30 ^d 9 ^d
Resident – Adult Resident – Child	30 ^d 9 ^d
Resident - Child	0 6
BW = Body weight (kg)	
Adult	70 ^d 70 ^d
Child	15 ^d 15 ^d
AT _{carc} = Averaging time for carcinogens (days)	25,550 ^d 25,550 ^d
AT _{non} = Averaging time for noncarcinogens (days	
<u> </u>	days/year days/year
F:\0	ATAUobs\Port of Portland\18191-01 T-1 Risk Assessmeni\Tables\TableSoll-Ing(T1)

Notes:

(a) Lifetime average daily dose, the intake value used to evaluate potential carcinogenic effects. For the residential evaluation, the adult and child intakes will be combined as recommended in Appendix A, Section A.O.of.DEQ guidance (2000).

(b) Average daily dose, the intake value used to evaluate potential noncarcinogenic effects.

(c) An upper one-sided 90 percent confidence limit of the mean or the maximum concentration (whichever is lower) used for the RME.

(d) DEQ (December 2000).

(e) Reasonable maximum exposure.

(1) Central Tendency.

Table 6 - Exposure Dose Equations and Exposure Factor Values: Dermal Contact with Soil

Marine Terminal 1 South Risk Assessment Portland, Oregon

LADD ^a (mg/kg-d)	=	C _{soil} x AF x SA x DAF x EF x ED x CF
		DIA/ v. A4

 ADD^b (mg/kg-d) $\simeq C_{sol} \times AF \times SA \times DAF \times EF \times ED \times CF$

						! —
			Exposure Factor (units)	RME ^e Value	CT ^f Value	
	C _{soli}	· -	Chemical concentration in soil (mg/kg)	UCL ₉₀	Arithmetic Mean	· ·
	AF	_	Soil-to-skin adherence factor (mg/cm²-event)			\vdash
	Δ-	-	Commercial Worker	0.08 ^d	0.08 ^d	
	٠.		Utility/Excavation Worker	1.0 ^d	0.3 ^d	ĺ
	 		Resident – Adult	0.08 ^d	0.08 ^d	├
			Resident - Child	1.0	0.08 0.3 ^d	├
	i		Nesidetit – Chiid	1.0	0.3	1
	ŞA	=	Skin surface area (cm²/day)			i
			Commercial Worker	4100 ^d	3200 ^d	
			Utility/Excavation Worker	4100 ^d	3200 ^d	
	İ		Resident - Adult	6900 ^d	5200 ^d	ĺ
	}		Resident – Child	5000 ^d	4500 ^d	İ
				}		İ
	DAF	=	Dermal absorption factor (unitless)	Chemical-specific	Chemical-specific	
	EF	=	Exposure frequency (days/year)	arad	250 ^d	
			Commercial Worker	250 ^d 9 ^d	25U 9 ^d	
,			Utility/Excavation Worker		40 ^d	-
			Resident – Adult/Child	350 ^d	40	H
		_	Financian disastina (see)			
	ED	=	Exposure duration (years)	ord	6 ^d	
			Commercial Worker	25 ^d	0.5 ^d	
			Utility/Excavation Worker Resident – Adult	30 ^d	0.5 9 ^d	
			Resident – Adult Resident – Child	50 6 ^d	9 6 ^d	İ
			Resident - Child	, b	O	İ
	-CF		-Conversion factor (kg/mg)	10-6	10-8	L
	OF .	-	- Conversion factor (kg/mg)	10	. 10	
	BW	=	Body weight (kg)	70 ^d	70 ^d	
			Adult	15 ^d	15 ^d	İ
			Child	1		Ì
	AT _{carc}	=	Averaging time for carcinogens (days)	25,550 ^d	25,550 ^d	
	AT _{non}	=	Averaging time for noncarcinogens (days)	ED (years) x 365	ED (years) x 365	
				days/year ^d	days/year ^d	

Notes:

(b) Absorbed daily dose, intake value used to evaluate potential noncarcinogenic effects.

(4) DEQ (December 2000).

(f) Central Tendency.

F:IDATA Jobs Port of Portland (15191-01.T-1. Risk Assessment Tables (Table 6Derm (T1)

⁽⁴⁾ Lifetime absorbed daily dose, intake value used to evaluate potential carcinogenic effects. For the residential evaluation, the adult and child intakes will be combined as recommended in Appendix A, Section A.0 of DEQ guidance (2000).

⁽c) An upper one-sided 90 percent confidence limit of the mean or the maximum concentration (whichever is tower) was used for the RME.

⁽e) Reasonable maximum exposure.

Table 7 - Exposure Dose Equations and Exposure Factor Values:

Inhalation of Volatiles

Marine Terminal 1 South Risk Assessment

Portland, Oregon

ADD ^b (mg/kg·d) = C _{sir,X} Rx EF x ED BW x At _{ron} Exposure Factor (units) RME ^f Value CT ^g Value Contained and a concentration in air (mg/m ^g) UCL ₉₀ Arithmetic Mean IR = Inhalation rate (m ^g /day) Commercial Worker 15.2 ^e 15.2 ^e Utility/Exeavation Worker 15.2 ^e 15.2 ^e Resident – Adult 15.2 ^e 15.2 ^e Resident – Child 8.3 ^e 8.3 ^e EF = Exposure frequency (days/year) Commercial Worker 9 ^e 9 ^e Resident – Adult/Child 350 ^e 350 ^e ED = Exposure duration (years) Commercial Worker 1 ^e Utility/Excavation Worker 1 ^e Resident – Adult 30 ^e Resident – Adult 30 ^e Resident – Adult 30 ^e Resident – Child 6 ^e Resident – Child 70 ^e Resident – Chi		$LADD'''(mg/kg-d)''' = \underline{C}_{alr} \underline{x}$	BW x At _{carc}	
Colf = Chemical concentration in air (mg/m³) IR = Inhalation rate (m³/day) Commercial Worker Utility/Exeavation Worker Resident – Adult Resident – Child EF = Exposure frequency (days/year) Commercial Worker Utility/Excavation Worker Po Po Po Po Po Po Po Po Po Po Po Po Po P	•	ADD ^b (mg/kg-d) = - <u>G_{air}x</u>	IR-x-EF-x-ED BW x At _{non}	
IR		Exposure Factor (units)	RME ^r Value	CT ^g Value
Commercial Worker		C _{eir} = Chemical concentration in air (mg/m³)	UCL ₈₀ °	Arithmetic Mean
Utility/Exeavation Worker 15.2° 15.2° Resident - Adult 15.2° 15.2° Resident - Child 8.3° 8.3° EF = Exposure frequency (days/year) Commercial Worker 250° 250° Utility/Excavation Worker 9° 9° Resident - Adult/Child 350° 350° ED = Exposure duration (years) Commercial Worker 25° 6° Utility/Excavation Worker 1° 0.5° Resident - Adult 30° 9° Resident - Child 6° 6° BW = Body weight (kg) Adult 70° 70°			·	
Resident - Adult 15.2° 8.3° 8.3° EF				15.2
Resident - Child 8.3° 8.3°			15.2°	15.2
Commercial Worker		· •	15.2° 8.3°	15.2° 8.3°
Commercial Worker		EF = Exposure frequency (days/year)		
Resident - Adult/Child 350° 350° ED = Exposure duration (years) Commercial Worker 25° 6° Utility/Excavation Worker 1° 0.5° Resident - Adult 30° 9° Resident - Child 6° 6° BW = Body weight (kg) Adult 70° 70°			250°	250°
ED = Exposure duration (years) Commercial Worker Utility/Excavation Worker Resident—Adult Resident—Child BW = Body weight (kg) Adult 70e Frequence of the commercial worker of the commercia				
Commercial Worker 25° 6° Utility/Excavation Worker 1° 0.5° Resident—Adult 30° 9° Resident—Child 6° 6° BW = Body weight (kg) 70° 70°		Resident - Adult/Child	350°	350°
Utility/Excavation Worker				- · · · · · · · · · · · · · · · · · · ·
Resident Adult 30e 9e 6e 6e 6e 6e 6e 6e 70e 70e 70e 70e			25°	
Resident—Child—6° 6° BW = Body weight (kg) Adult 70° 70°				
BW = Body weight (kg) Adult 70° 70°				9 9
Adult 70° 70°		Kesident - Child	69	6
Adult 70 ^e 70 ^o		BW = Body weight (kg)		
Child 15 ⁶ 15 ⁰		Adult		
	•	Child	15 ^e	15°

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25,550^e

ED (years) x 365

days/year-

25,550°

ED (years) x 365

-days/year

(b) Average daily dose, intake value used to evaluate potential noncarcinogenic effects.

AT_{care} = Averaging time for carcinogens (days)

Averaging-time-for-noncarcinogens

(days)

(9) Central Tendency.

⁽a) Lifetime average daily dose, intake value used to evaluate potential carcinogenic effects. For the residential evaluation, the adult and child intakes will be combined as recommended in Appendix A, Section A.0 of DEQ guidance (2000).

⁽c) Upper one-sided 90 percent confidence limit of the mean or the maximum concentration (whichever is lower) was used for the RME.

(d) C_{str} was derived from soil and groundwater concentrations using models discussed in DEQ guidance (1999 and 2000).

(e) DEQ (December 2000).

n Reasonable maximum exposure

9e

350^e

Table 8 - Exposure Dose Equations and Exposure Factor Values: Inhalation of Dust Marine-Terminal-1-South-Risk-Assessment-

Portland, Oregon

Utility/Excavation Worker

Resident - Adult/Child

Exposure duration (years)

_	LADD ^a (mg/kg-d) = PM ₁₀ x II	R-x-EF-x-ED W x At _{eare}		_
	ADD ^b (mg/kg-d) = PM ₁₀ .x-li	R x EF x ED V x At _{ron}		
_		Y A CUROR		_
	Exposure Factor (units)	RME ^f Value	CT ^g Value	
_	PM ₁₀ ^d = Respirable particulate concentration in air (mg/m³)	UCL ₉₀	Arithmetic Mean	_
_	IR = Inhalation-rate (m³/day) Commercial Worker	15.2 ^e	15.2 ^e	_
	Utility/Excavation Worker Resident – Adult Resident – Child	15.2° 15.2° 15.2° 8.3°	15.2 ^e 15.2 ^e 15.2 ^e 8.3 ^e	
	EF = Exposure frequency (days/year) Commercial Worker	250°	250 ^e	_

96

350°

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į	Commercial Worker	25°	6 ^e
	Utility/Excavation Worker	18	-0.5°
Ī	Resident – Adult Resident – Child	30° 6°	6°
	BW = Body weight (kg)		

Adult 70° 70^e Child 15° 15^e 25,550° Averaging time for carcinogens (days) 25,550e

Averaging time for noncarcinogens ED (years) x 365 ED (years) x 365 (days) days/year days/year

Notes:

(a) Lifetime average daily dose, intake value used to evaluate potential carcinogenic effects. For the residential evaluation, the adult and child intakes will be combined as recommended in Appendix A, Section A.0 of DEQ guidance (2000)."

(h) Averago dally dose, intake value used to evaluate potential noncarcinogenic effects.

(c) Upper one-sided 90 percent confidence limit of the mean or the maximum concentration (whichever is lower) was used for the RME.

(4) PM₁₀ was derived using the Particulate Emission Factor equation presented in DEQ guidance (2000).

(A) DEQ (December 2000).

(1) Reasonable maximum exposure.

(9) Central Tendency.

ED

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Table 11 - RME Risk Summary: By COPC
Marine Terminal 1 South Risk Assessment
Portland, Oregon

					F	ME Cancer	Risk		ĺ
	0.4	F 0	0000	1 15		Inhalation	Inhalation		1
	SubArea	Exposure Scenario	COPC	Ingestion	Dermal	of Volatiles	of Dust	TOTAL	
								THE POST	
	Area A	Resident	Benzo(a)anthracene	4.E-06	6.E-06	na	1.E-10	S(1505)	
=	1	Ì	Benzo(a)pyrene	4.E-05	5.E-05	na	1.E-09	9E054	
			Benzo(b)fluoranthene	3.E-06	4.E-06_	na	8.E-11	e de ida	
			Dibenz(a,h)anthracene	3.E-06	5.E-06	na	1.E-10_	4 /8 (E 4 / 6 · 2	
1000 ·	1		Indeno(1,2,3-cd)pyrene	1.E-06	2.E-06	na	4.E-11		l
	1	ĺ	Arsenic	3.E-05	1.E-05	na	2.E-08	÷5 € 05	1
	1		Tetrachloroethene	na	na	4.E-09	na	4.E-09	i
		·	TOTAL	8.E-05	.8.E-05	4.E-09	3.E-08	2.E-04	
٨									
. 🖷	1	Commercial Worker	Benzo(a)anthracene	5.E-07	2.E-07	na	3.E-11	7.E-07	l
	1		Benzo(a)pyrene	5.E-06	2.E-06	na	2.E-10	7.E-06	
			Benzo(b)fluoranthene	4.E-07	2.E-07	na	-2.E-11	5.E-07	
			Dibenz(a,h)anthracene	4.E-07	2.E-07	na	2.E-11	6.E-07	
 -			Indeno(1,2,3-cd)pyrene	2.E-07	8.E-08	na ,	9.E-12	3.E-07	ł
-	1		Arsenic	4.E-06	4.E-07	na	5.E-09	45 E1061V	-
	†		Tetrachloroethene	na	na	6.E-10	na	6.E-10	
	 		TOTAL	1:E:05	3.E-06	6.E-10	5.E-09	1.E-05	
	1								
	Area B	Resident	Benzo(a)anthracene	3.E-07	4.E-07	na	9.E-12	7.E-07	
	1		Benzo(a)pyrene	4.E-06	5.E-06	na	1.E-10	91706	
	+		Benzo(b)fluoranthene	3.E-07	4.E-07	na	8.E-12	6.E-07	
.			Indeno(1,2,3-cd)pyrene	2.E-07	3.E-07	na	7.E-12	6.E-07	
	1		Arsenic	1.E-05	4.E-06	na	9.E-09	ZE05 2	
. 	†		TOTAL	2.E-05	1.E-05	na	9:E-09	3.E-05	
	1.		· · · · · · · · · · · · · · · · · · ·						
		Commercial Worker	Benzo(a)anthracene	4.E-08	2.E-08	na	2,E-12	5.E-08	
•			Benzo(a)pyrene	5.E-07	2.E-07	· na ·	2.E-11	7.E-07	
 -	 		Benzo(b)fluoranthene	3.E-08	-1.E-08	na	2.E-12	-5.E-08	-
	-		Indeno(1,2,3-cd)pyrene	3.E-08	1.E-08	na	2.E-12	4.E=08	 -
	}		Arsenic	2.E-06	2.E-07	na	2.E-09	22 E 06 E	
	}		TOTAL	2.E-06	4.E-07	na	2.E-09	3.E-06	
									
	- Area C	Resident	Arsenic	1.E-05	4.E-06	na ,	8:E-09	2/E-05	
					ĺ	· J			l
=		Commercial Worker	Arsenic	2.E-06	2.E-07	na	2.E-09	2 2 € 06	
				:DATAUJobsvPort o	Fortland\15191	O1 T-1 Risk Assessn			
	Mada								

Note:

^{1.} Shaded boxes indicate COPC that exceeds DEQ acceptable risk target.

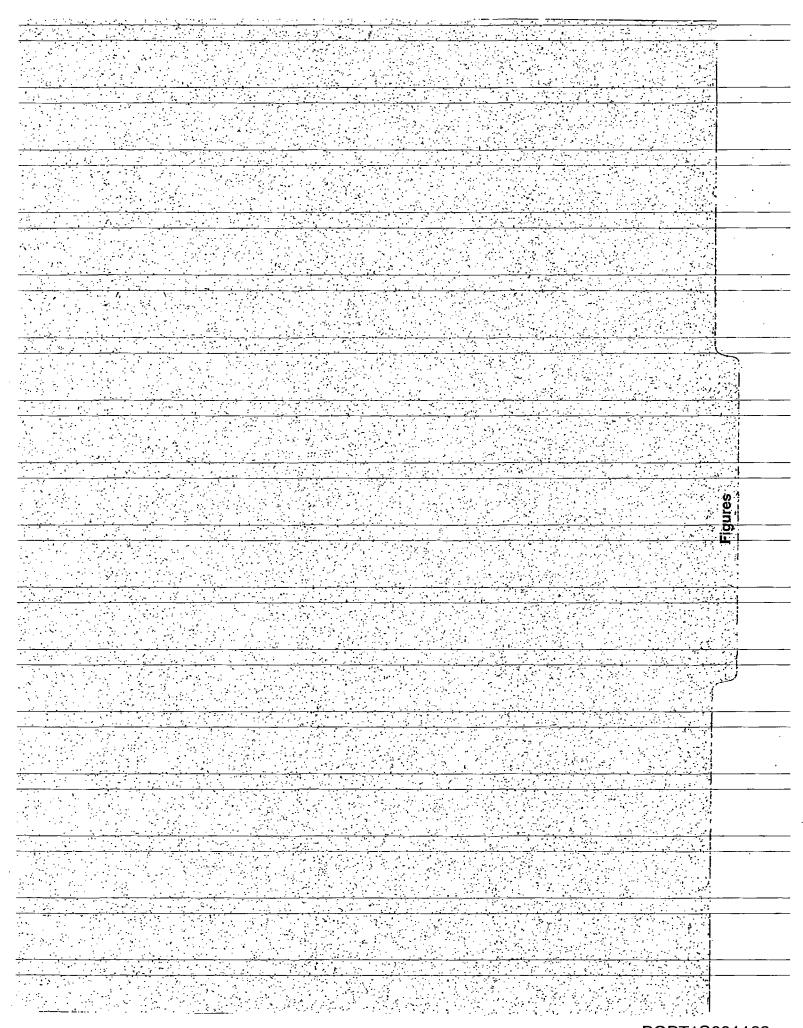
Table 12 - Modified Level 2 Screening of Groundwater Results Marine Terminal 1 South Risk Assessment Portland, Oregon

Sheet 1 of 1

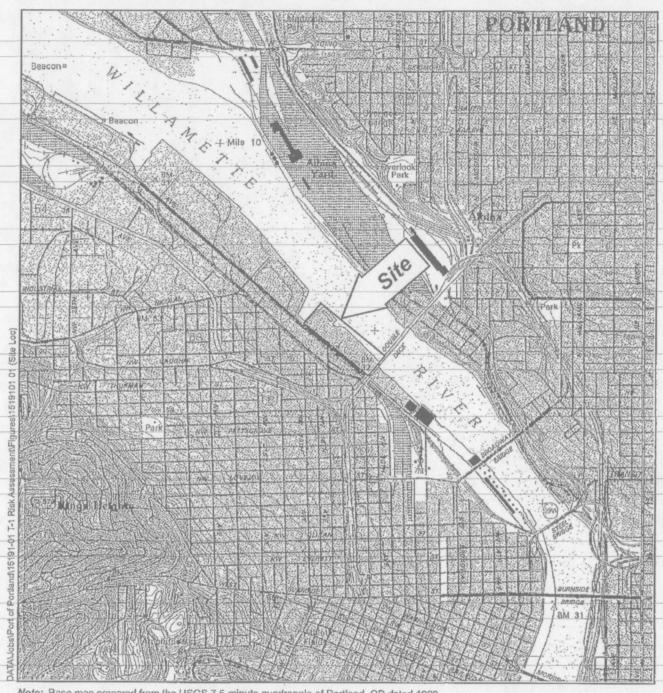
Sampling Date	10/01/2001	**************************************							
		(01011200)	9/28/2001	-9/28/2001	9/28/2001	9/28/2001	10/01/2001	Aquatic	
Total Metals in µg/L									
Arsenic	2.01	12.8	14	6.45	12.1	2.72	1.38	150	
Cadmium									
		1 11						_	
		NΔ	NA.						,
	1.11								
	10.8								_
		- 101		0.00			11.0	120	
		14.5	11	6.51	11.3	3.65	1 11	150	
	2.29								
Lead	1.37								-
PAHs in pg/L									
	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U	0.1 เป	0.027	
	0.1 U	0.1 U							
Benzo(b)ffuoranthene	0.1 U	0.1 U			0.1 U			_	·
Benzo(k)fluoranthene	0.1 U	0.1 U			0.1 U				
Chrysene	0.1 U	0.1 ป			0.1 U				
Dibenz(ah)anthracene	0.2 U	0.2 U							
Indeno(1,2,3-cd)pyrene	0.1-U-	0,1 U							
Acenaphthene	0.1 U	0.121	0.192					520	
Acensphihylene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U			_	
Anthracene	0.1 U	0.1 U	0.1 ป	0.1 U	0.1 U	0.1 U	0.1 U	13	
Benzo(ghi)perylene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U		
Fluoranthene	0.1 ป	0.119	0.1 U	0.1 U	0.1 ป	0.1 U	0.1 U	6.2	·
Fluorene	0.1 U	0.1 U	0.1 U	0.10	0.1 U	ט־1.10	0.1 U	3.9	
Naphthalene	0.1 U	0.1 U	0.1 U	0.291	0.1 U	ט 1.0	0.1 U	620	
Phenanthrene	0.1 ป	1.25	0.138	0,576	1.18	0.1 U	0.153	6.3	
Pyrene	0.1 U	0.564	0.1 U	0.123	0.172	0.1 U	0.153	_	
Total PAHs	0.2-U	2.051	0:33	1.71	1 . 78	- 0.2 U			
Volatiles in µg/L									
Benzene .		NA	· NA	1 0	NA	NA .	NA	130	
DEHP	10 U	10 U	NA NA	NA.	NA	NΑ	10 U	3	•
Ethytbenzone	1 U	NA	NA	10	NA.	NA	NA	7.3	
1		NA	NA	10	NA	NA	NA	840	•
Toloene	10	NA	NA.	10	NA NA	NA.	NA ·	9.8	
Total xylenes	1 U	NA	NA.	1_U.	. NA	NV	NA	13	·
	Chromium Copper Lead Marcury Nickel Silver Zinc Disadved Motals in µg/L Arsenic Copper Lead PAHs in µg/L—Benzo(s)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Benzo(b)fluoranthene Chrysene Dibenz(ah)anthracene Indene(1,2,3-ed)pyrene Acenaphthene Acenaphthylene Anthracene Benzo(ghi)perylene Fluoranthene Fluoranthene Fluoranthene Phenanthrene Phenanthrene Pyrene Total PAHs Volatiles in µg/L Benzene DEHP Ethylbenzene PCE	Chromium 3.25	Chromium 3.25	Chromium 3.25	Chromitum 3.25	Chromium 3.25	Chromium 3.25	Chromium 3.25	Chromium 3.25

Notes:

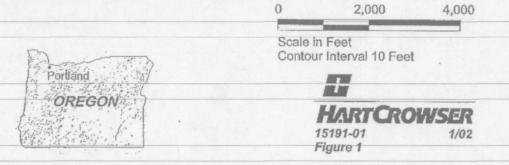
- 1. U = Not Detected at Reported Detection Limit.
 2. Na = Not Analyzed.
 3. Shading Denotes Analyte Exceeding Ecological Screening Benchmark Value.
 4. DEQ Ecological Screening Benchmark Value Freshwater Value for Aquatic Organisms (DEQ March 2001).

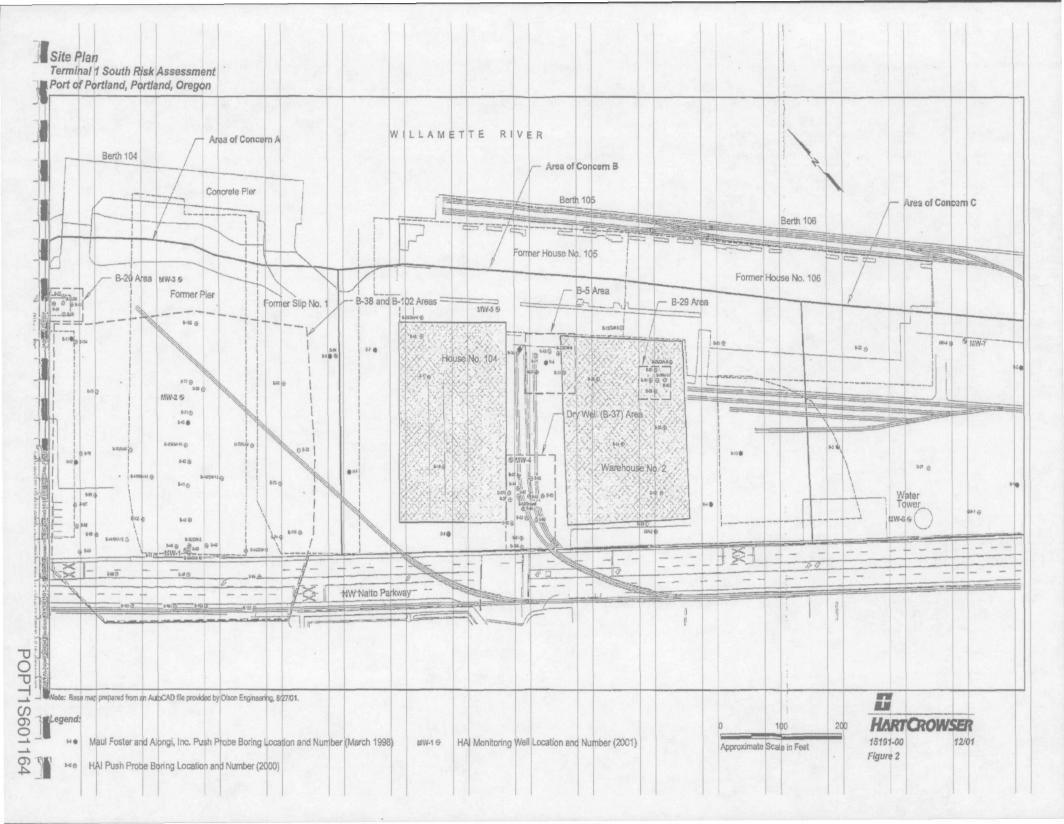


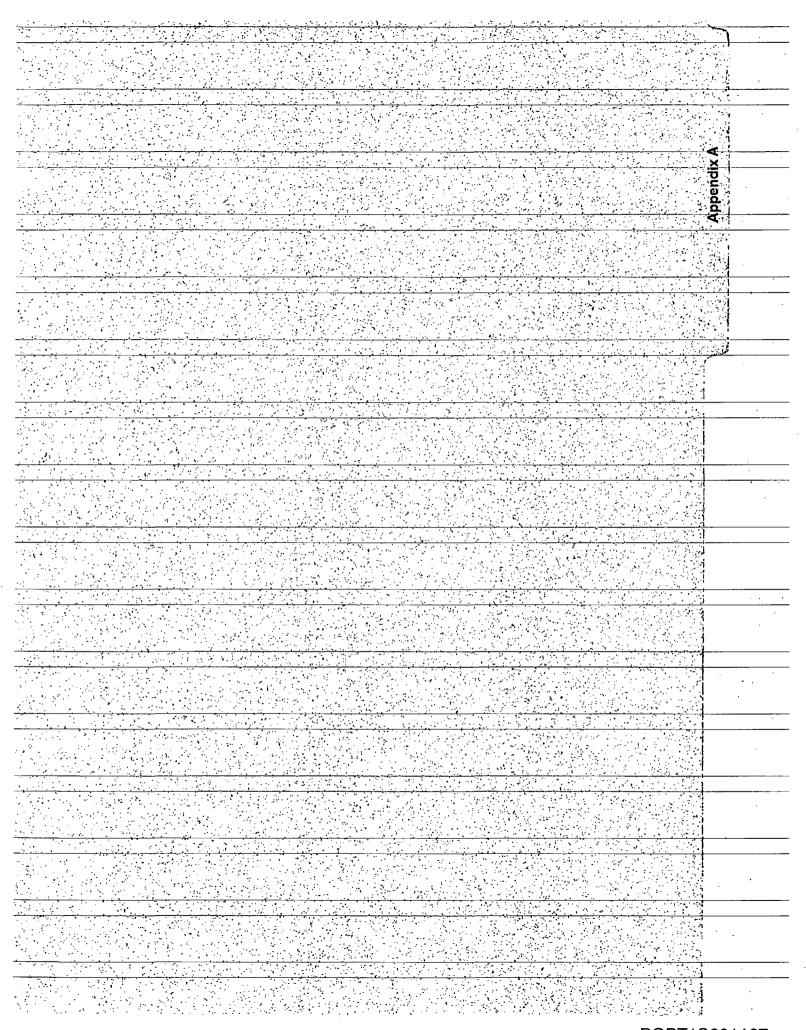
Site Location Map Terminal 1 South Risk Assessment Port of Portland, Portland, Oregon



Note: Base map prepared from the USGS 7.5-minute quadrangle of Portland, OR dated 1990.







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			Recorded States		
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Hart Crowser					
Hart Crowser 15191-01 January (8, 2002					
		SOIL AND GROUNDWA	SOIL AND GROUNDWATER ANALY	SOIL AND GROUNDWATER ANALYTICAL DATA US RISK-AS	SOIL AND GROUNDWATER ANALYTICAL DATA USED FOR THE RISK ASSESSMENTS:

Station Sampling Date Depth in Feet	B-100	B-100) B-10	6/2000	5105-001025-060 B-102 10/26/2000 10	5106-001027-074 B-103 10/27/2000 10	5106-001030-079 B-104 10/30/2000 2.5	5105-001027-078 B-105 10/27/2000 2.5	5106-01027-067 B-107 10/27/2000 4	5105-001026-057 B-108 10/26/2000 2.5	B11 (9-11) B-11 3/28/98 9-1	5108-001026-056 B-110 10/26/2000 2.5	B-38 3/01/2000 10	B 40 3/13/2000 5
Metals in mg/kg Animony Arsenic							1.64	1 U	1.35	1.49	0.5.UJ 11.2 J	U		
Beryllium Cadmium	<i>i</i>										0.5 U ∎.33 28.4		02 U 223	
Chromium Copper Lead							28.1	k u	2.73	17.1	17.1 23.8	.	807	
Mercury Nickel											0.05 U 15.2			
Selenium Silver											0.5 U			
Thaillum Zinc TCLP-Lead											0.5 U 70.5			
PAHs in mg/kg Benzo(a)anthr	ecans 0	05 U 0	.05 U	0.05 U	0.05 U	0.05 U			0.05 U	0,09	•	1 U	0.472 J	0.076
Benzo(a)pyrer Benzo(b)fluora	e 0.	อธบ 0	.05 U	0.05 U	0.05 U 0.05 U	0.05 U 0.05 U			0.05 U 0.055	0.135)).	10	0.743 J 1.57 J	0.086 0.096
Benzo(k)fluora Chrysene	inthene 0.	.05 U O	.05 U	0.05 U 0.05 U	0.05 U 0.05 U	0.05 U 0.05 U			0.05 U 0.05 U	0.055 0.075	-	1 U	0.01 UJ 2.18 J	0.025 0.102
Dibenz(ah)ant Indeno(1,2,3-c	d)pyrene 0	.05 U 0	.05 U	0.05 U 0.05 U	0.05 U 0.05 U	0.05 U 0.05 U			0.05 U 0.05 U	0.05 U 0.105		1 U	0.35 J 0.676 J	0.015 0.052
Acenaphthene Acenaphthyler	ne 0	.05 U 0	.05 U	0.05 U 0.05 U	0.05 U 0.05 U	0.05 U 0.05 U 0.05 U			0.05 U 0.05 U 0.05 U	0 05 U 0 05 U 0 05 U		. 1 U 1 U	0.63 J 0.35 J 0.883 J	0.022 0.021 0.028
Anthradene Benzo(ghi)per Fluorenthene	ylene 0	.05 U 0	.05 U	0.05 U 0.05 U 0.05 U	0.06 0.05 U 0.05 U	0.05 U			0.05 U 0.09	0.165		1 0	1.28 J 0.938 J	0.067 0.139
Fluorena Naphthalena	0	.05 ປ 0	.05 U	0.05 U 0.05 U	0.05 U 0.05 U	0.05 U 0.05 U			0.05 U 0.05 U	0.05 U 0.05 U		1 U	0.928 J 0.957 J	0.013 0.013
Phenarthrene Pyrene	0	.05 ป 0	.05 U	0.05 U 0.05 U	0.05 U 0.075	0.05 U 0.05 U			0.05 U 0.05 U	0.165 0.37		1 U	2.35 J 2.83 J	0.104 0.209
Total PAHs	•			0.05 U	0.135	0.05			0.145	1,61		1 U	17.132	1.068
	Plea	se refer to notes	at end of table.											
							. .							
\ \													.	.

Depth in Feet Motals in mg/kg	Results for 3 A 4876-000313-043 B-41 3/13/2000 5	Soil Samples A 4876-000313-056 B-44 3/13/2000	(Metals and P, A 4878-000316-087 B-46 3/16/2000 12.5	AHs) A 4876-000316-071 B-47 3/16/2000 2.5	A 4876-000316-0 B-48 3/16/2000 11	A 79 5105-000921-044 B-53 9/21/2000	A 5106-000921-050 B-68 9/21/2000 10	A 8-68-Duplicate 8-68 9/21/2000 10	A 5105-000321-049 B-68 9/21/2000 2.5	A 5106-000921-053 B-69 9/21/2000 10	A 5106-000921-052 B-69 9/21/2000 2.5	A 5106-0003921-057 B-70 9/21/2000 10	Sheet 2 of 6 A 5106-00921-058 B-70 9/21/2000 2.5
Antimony Arsenic Betylllum Cadmium I Chromium Copper Lead Mercury Nicket Selenium Silver Thalium Zinc						0.5 U 1.82 0.5 U 0.5 U 14.8 13.3 10 U 0.1 U 13.7 0.5 U 0.5 U			19.5 12.9 0.5 U 7,43 48.2 286 6190 9,68 64.3 0.5 U 1 U 3.5		1 d U		
TC_P-lead PARs in mg/kg Behzo(a)anthracene Behzo(b)pyrene Benzo(b)fluoranthene Benzo(k)fluoranthene Chrysene Ditenz(ah)anthracene Indeno(1,2,3-cd)pyrene Acenaphthene Acenaphthylene Anthracene Benzo(ghi)perylene Fluoranthene Fluoranthene Phenanthrene Phenanthrene Pyrene	0.062 0.068 0.069 0.019 0.08 0.015 U 0.034 0.053 0.015 U 0.031 0.049 0.169 0.044 0.034 0.178 0.195	0.01 U 0.01 U 0.018 0.01 U 0.021 0.01 U 0.01 U 0.01 U 0.01 U 0.01 U 0.01 U 0.01 U 0.01 U 0.015 0.015	0.074 0.096 0.084 0.023 0.073 0.014 U 0.042 0.014 U 0.015 0.019 0.058 0.101 0.014 U 0.026 0.048	0.227 0.243 0.242 0.082 0.258 0.033 0.113 0.026 0.046 0.067 0.16 0.371 0.034 0.082 0.235 0.235	0.135 R. R. A. A. A. A. A. A. A. A. A. A. A. A. A.	0.444 0.419 0.224 0.349 0.55 0.0894 0.233 0.0268 U 0.114 0.117 0.27 0.438 0.0268 U 0.0268 U	0.139 0.134 0.131 0.104 0.18 0.0268 U 0.0873 0.0268 U 0.0528 0.115 0.197 0.0268 U 0.0268 U 0.0528	0.139 0.134 0.181 0.104 0.18 0.0268 U 0.039 0.0268 U 0.0268 U 0.115 0.102 0.0268 U 0.0268 U	295 662 935 705 422 553 956 134 U 338 658 114 376 195 572 7.9 34.8 27.6	0.0268 U 0.0268 U 0.0268 U 0.0309 0.0268 U 0.0268 U 0.0268 U 0.0268 U 0.0268 U 0.0268 U 0.0268 U 0.0268 U 0.0268 U	0.0134 U 0.0134 U	0.11 9.118 9.063 9.083 9.129 9.0176 9.024 9.024 9.0552 9.0563 9.111 9.204 9.0167 9.0167 9.0151 9.267 9.416	0.0268 U 0.0268 U
Total PAHs	1.085	0,154 notes at end of	0.81	2.755	1,994 R	4.2514	1.5616	1.1083	156 35	0.1431	0.0134 U	1.7677	C 0248 U

B B B B B B B B B B B B B B B B B B B	0.5 U 0.5 U 39 3 0.132 0.167 0.1 0.101 0.147 0.067 U 0.067 U 0.067 U 0.067 U 0.067 U 0.067 U 0.126 0.285 1.4741	
B	0.132 0.167 0.1 0.101 0.147 0.067 0.0891 0.067 0.067 0.118 0.209 0.067 0.067 0.067 0.126 0.285	
B	0.132 0.167 0.1 0.101 0.147 0.067 U 0.0891 0.067 U 0.067 U 0.118 0.209 0.067 U 0.126 0.285	
5105-001024-008	0.132 0.167 0.1 0.101 0.147 0.067 0.0891 0.067 0.067 0.118 0.209 0.067 0.067 0.067 0.067 0.126 0.285	
5105-001024-008	0.132 0.167 0.1 0.101 0.147 0.067 0.0891 0.067 U 0.067 U 0.118 0.209 0.067 U 0.126 0.285	
5106-001024-008 4876-0 B-106 B-14 10/24/2000 2/29/ 7 1	0.132 0.167 0.1 0.101 0.147 0.087 J 0.087 J 0.067 J 0.067 J 0.067 J 0.067 J 0.118 0.209 0.067 J 0.126 0.285	
5106-001024-008 B-106, 10/24/2000	0.167 0.1 0.101 0.147 0.067 0.067 0.067 0.067 0.118 0.209 0.067 0.067 0.067 0.067 0.067 0.067 0.067 0.067	
5106-00 B-106, 10/24/2	0.16 0.10 0.14 0.06 0.089 0.06 0.06 0.06 0.06 0.12	
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A 5105-00 B-99 11/06/2 3		
05-001106-09 99 1/06/2000	0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U	
E		
A 5106-00110 B-97 11/06/200 5	0.05 0.05 0.05 0.05 0.05 0.05 0.05 0.05	
7:53	0.05 U U 0.05 U 0.05 U	
-397 S		
A 5105-001108 B-97 11/06/2000	0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U 0.05 U	
Hs) A 5106-001030-081 B-94 10/30/2000 2.5	1.26 1.36 1.7 0.57 1.31 0.16 0.55 0.19 0.12 0.55 0.79 2.22 0.135 0.13 1.9 2.86 15.805	
30-082	ט טטט טט	
(Metals an A 5106-001030 B-94 10/30/200 10	0.13 0.185 0.21 0.065 0.13 0.05 t 0.05 t 0.05 t 0.165 0.295 0.5 t 0.13 0.35 1.77	IZIOIE,
000921-070	067 U 067 U 067 U 067 U 067 U 067 U 067 U 067 U 067 U 067 U 067 U 067 U 067 U	at end of
71 510 B-f		
A 5108-000921-0 B-82 9/22/2000 10 0.532 1.76 0.5 U 0.5 U 12.5 12 10 U 1.53 J 14.4 0.5 U 1 U	0.5 U 45.1 0.067 U 0.067 U	Please refe
g Date Feet mg/kg y	and ng/kg)anthracerre)pyrene)fluoranthene e eh)anthracerre 1,2,3-cd)pyrene ithene ene hilpenytene hilpenytene hene elene hrene	•
Area Sample I Station Sampling Debth in Metals in Antimom Arsenid Betyllium Cadmiur Cadmiur Copper Lead Mercury Nickel Seleniun Silver	Benzo(a Benzo(k Benzo(k Chrysen Dibenz(a Indeno(Acenap Acenap Anthrace	
		DOT 1 2001 170

Station Sampling Date Depth in Feet Metals in mg/kg	B-52 9/19/2000 2	B-58	0.27/22-		5106-001024-037		5106-00/1024/043	5106-001024-049	5106-001024-047	5106 001080-085	5106-001030-085	5106-00 024-032	
		9/19/2000	8-63/63a 9/20/2000	B-64/64a 10/25/2000 10	B-64/64a 10/25/2000 2.5	B-65/6\$a 10/25/2000 12	B-65/66a 10/25/2000 2.5	8-66/66a 10/25/2000 10	B-66/66a 10/25/2000 2.5	B-67/674 10/30/2000 13.5	B-67/67a 10/30/2000 5.5	B-87 10/25/2000 2.5	B-88 10/25/20 6.5
Metals in mg/kg		•	10.5		2.5	12	2.0	10	2.5	13.3	3.5	2.3	<u>"</u>
Antimony	0.5 บ	0.5 U	0.703					•					
Arsenic	2.6	2.05	3.6		1						1 1		
Beryllum	0.5 U	0.5 U	0.5								1		
Cadmium	1.34	0.584	0.703		.								
Chromium	16.4	11.2	6.97		1 1								[]
Copper	27.1	11	16,4	r ·	.] [•		.		
Lead	121	10 U	92									.	
Mercury	0.128	0.1 U	0.145					·			}	-	
Nickel Selenium	23.5	16	12	ľ. I I		·	·]				.	
Silver	0.0613 1 U	0.5 U 1 U	0.5 U					!		.			
Thellium	0.5 U	0.5 U	0.5 U	i i.l.	•						•	- 1	
Zinc	140	41.5	107						.				
TCLP-Lead		1			.								
PAHs in mg/kg								- 1 1					
Benzo(a)anthracene	0.0448	0.02	67 U	0.403	0.0524	1.68 U	0.087 U	1.18	0. 067 U	0.825	0.05 U	0.149	1.\$1
· Benzo(a)pyrene	0.0501	0.0246	67 U	0.605	0.0834	1.68 U	0.067 U	1.16	0. 0 67 U	0.92	0.05 U	0.188	1.45
Benzo(b)fluoranthene	0.0402	0.0215	67 U	0.4	0.134	1.68 U	0.067 U	1.28	0. 0 67 U	0.82	0.05 U	0.116	0.902
Benzo(k)fluoranthene	0.0471	0.0177	67 U	. 0.334	0.0934	1.68 U	0.067 U	0.844	0. 0 67 U	0.315	0.05 U	0.114	0.919
Chrysene	0.0666 0.0268 U	0.0229	67 U 67 U	0.465 0.0839	0.0798	1.88	0.067 U 0.067 U	1.58 0.22	0.067 U 0.067 U	0.67 0.05 U	0.05 U 0.05 U	0.163 0.067 U	1.57 0.247
Dibenz(ah)anihracene Indeno(1,2,3-od)pyreno	1	0.0134 U	67 U	0.0839	0.0268 U 0.117	1.68 U 1.68 U	0.067 U	0.748	0.067 U	0.325	0.05 U	0.102	0.69
Acenaphthene	0.0268 U	0.0134 U	106	0,0754	0.0268 U	1.68 U	0.067 U	0.516	0.067 U	0.05 บ	0.05 U	0.067 U	0.067
Acenaphthylene	0.0314	0.0134 U	67 U	0.0765	0.0268 U	1.68 Ü	0.067 U	0.263	0. 0 67 U	0.05 U	0.05 U	0.067 U	0.067
Arthracene	0.0268 U	0.0134 U	67.8	0.136	0.0268 ป	1.68 U	0.067 U	0.464	0. 0 67 U	0.185	0.05 じ	0.067 U	0.156
Benzo(ghi)perylene	0.0536	0.0262	67 U	0.564	0.174	1.68 U	0.067 U	0.837	0.067 U	0.405	0.0 5 U	0.136	0.88
· Fluoranthene	0.0977	0.037	285	1 1	0.0904	1,68 U	0.067 U	4.5	0.067 U	0.72	0.05 U	0.219	1.5
Fluorene	0.0268 U	0.0134 ป	174	0.067 U	0.0268 U	1.68 U	0.067 U	0.974	0.067 U	0.05 U	0.05 U	0.067 U	0.067
Naphthalene	0.0268 U	0.0134 U	67 U	0.104	0.0268 U	1.68 U	0.067 U	3.99	0. 0 67 U	0.05 U	0.05 U	0.067 U	0.067
Phenanthrene	0.0588	0.0164	700	0.583	0.0395	1.68 U	0.067 U	3.73	0.067 U 0.067 U	0.34 1.04	0.05 U 0.05 U	0.137 0.278	0.77 2.74
Pyrene Total PAHs	0.0941	0.0314 0.237	143 1475.8	1.36 6.5738	. 0.0997 0.9636	1.68 U 1.88	0.067 U 0.067 U	3.97 26.226	0.067 U	6.565	0.05 U	1.502	13.334
							1.5						
	Pibase leter (o notes at end of	table.								. .		-
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12.4 9 14.4	
19 U 165 2.5 U 0.65 U 0.7 U 19.3 U 0.5 U	ble A-1 - Analytica Area Sample ID Station Sampling Date Depth in Feet Stats in mg/kg Arsenic Ben/lium Cadmium Chromium
0.64 U 0.65 U 14 U 0.409 1.32 0.5 U 0.5 U 1 U 0.5 U 0.	Results for 8 5105-001024-021 B-89 10724/2000 2.5
10 U 155 2,5 U 0,6 U 0,7 U 0,5 U 0,7 U 0,5	Soil Samples B 5106-001024-017 B-90 10/24/2000
19 U 16,5 U 0,5 U 0,5 U 1 U 0,3 U 0,5 U 1,5 U 0,5 U 1,5 U 0,5 U 1,5 U 0,	8
1.32 2.35 1.54 1.09 1.62 0.253 1.56 0.249 0.158 0.331 2.48 4.22 0.134 0.246 1.84 6.69 26.091	В
10 U 15,5 2,5 U 0.05 U 0.1 U 18,9 17,8 14.1 U 1 U 0.3 U 0.5	B 5108-001024-013 B-92 10/24/2000 10
15,5	C B2 (9-11) B-2 3/26/98 9-11 0.5 UJ 2.72 J 0.5 U 0.666 18.1
2.5 U 0.1 U 14.1 1 U 0.3 U 0.5 U 38	C B3 (11-12) B-3 3/25/94 11-12 0,5 UJ 11,8 J 0,782 1,3 25,7 19
	C 4876-00302-026 B-32 3/02/2000 1 2.5 UJ 2.9 0.18 0.2 U 11 14.4
	Sheet 6 of 6

iabic	A-2 - Analytical R	esuits for s	son Sampi	es (D	nesei anu C)(t)	Sheet 1 of 4
Area	Sample	Station	Sample	•		TPH-D in mg/kg	
	Number		Date	in fee	t Diesel	Oil	
Α	5106-001027-069	B-100	10/27/00	5	27 U	67.6 U	
	5106-001027-070	B-100	10/27/00	10	26.7 U	66.7 U	
A	5106-001027-071	B-100	10/27/00	20	29.9 U	74.6 U	
_	5106-001027-071	B-100 B-100	10/27/00	26	29.9 U	74.6 U	
. A	5106-001027-072	B-100 B-101	10/27/00	5	26.3 U	65.8 U	
A	5106-001026-064	B-101	10/26/00_		20.3 U	69.4 U	
	5106-001026-065	B-101	10/26/00- 10/26/00		276_U 28.2 U	70.4 U	
A	5106-001026-066	B-101	10/26/00	20 26	26.2 U 27 U	67.6 U	
A				5			
Α	5106-001026-059	—B-102——	10/26/00-		26.7-U	66.7-U	
— A	5106-001026-060 —	—B-102—	10/26/00	10	1170	1760	
A	5106-001026-061	B-102	10/26/00	20	28.2 U	70.4 U	
A	5106-001027-073	B-103	10/27/00	5	26.3 U	65.8 U	
A	5106-001027-074	B-103	10/27/00	10	27 U	67.6 U	
A	5106-001027-075	B-103	10/27/00	20	28.6 U	71.4 U	
Α -	5106-001027-076	B-103	10/27/00	26	28.6 U	71.4 U	
A	4876-000302-024	B-20	3/2/00	1	36	50 U	·
Α	4876-000301-020	B-38	3/1/00	10	34000 R	2500 UR - 3	
Α	4876-000301-021	B-38	3/1/00	20	500 R	50 UR	
A.	4876-000301-023	B-38	3/1/00	26	2.170 R	\$2,50 UR (
A	4876-000313-031	B-39	3/13/00	10	-25000 R		
A	4876-000313-036	B-39	3/13/00_	<u> 28</u>	æ #25 UR	56-50 UR-	
Α	4876-000313-037	B-40	3/13/00	5	910	50 U	
Α	4876-000313-038	B-40	3/13/00	10	25 U	50 U	
A	4876-000313-043	B-41	3/13/00_	5_	500	50 U	
A	<u> 4876-000313-044</u>	B-41	3/13/00 <u></u>	10	25 U	50 U	
Α	4876-000313-049	B-43	3/13/00	10	25 U	50 U	
Α	4876-000313-051	B-43	3/13/00	20	25 UR	7240 R	•
Α	4876-000313-053	B-43	3/13/00	27	25-U	50-U	
A	4876-000313-056	B-44	3/13/00	10	25 UR	230 R	
Α	4876-000313-058	B-44	3/13/00	20	25 U	50 U	
Α	4876-000313-060	B-44	3/13/00	26	25 Ú	50 U	
Α	4876-000316-062	B-45	3/16/00	10	25 U	50 U	
Α	4876-000316-063	B-45	3/16/00	18	360	50 U	
A	4876-000316-064	B-45	3/16/00	22	25 U	50 U	
Α	4876-000316-067	B-46	3/16/00	12.5	25 U	50 U	
Α	4876-000316-068	B-46	3/16/00	19	25 U	50 U	•
Α	4876-000316-069	B-46	3/16/00	25	25 U	50 U	
Α	4876-000316-071	B-47	3/16/00	2.5	450	50 U	
A	4876-000316-073	B-47	3/16/00	_10_	25 U	62	
A	4876-000316-075	B-47	3/16/00_	_20_	25_U	58	
Α	4876-000316-076	B-47	3/16/00	25	25 U	50 U	
A	4876-000316-078	B-48	3/16/00	6	25 U	160	
- A	4876-000316-079	B-48	3/16/00	-11-	25 U	1600	
— A	4876- 000316-080	B=48	3/16/00	15	25 U	77-	
A	4876-000316-082	B-49	3/16/00	5	254UR	50 UR	

Area	Sample	Station				TPH-D in mg/kg	
Alca	Number	Station	Date	in feet		Oil	
•	1070 000010 001	5.40	011000				
<u>A</u>	4876-000316-084	B-49	3/16/00		25 UR	A CONTRACTOR OF THE PROPERTY OF THE PARTY OF	
A	5106-000921-045	B-53	9/21/00	4	25 U	50 U	
Α	5106-000921-046	B-54	9/21/00	1	25 U	50 U	
A	5106-000921-043	B-55	9/21/00	1	25 U	50 U	
A	5106-000921-048	B-56	9/2 1/00-	1_	45.2	72.1	
A	5106-000921-047	B-57	9/21/00	1_	41.6		
Α	5106-000921-049	B-68	9/21/00	2.5	653	1130	
Α	5106-000921-050	B-68	9/21/00	10	25.5	84.3	
—A—	5106-000921-051	B-68	9/21/00	13-	199	386	
—A	B-68-Duplicate	B-68	9/21/00	10	32.4	84.9	······
Α	5106-000921-052	B-69	9/21/00	2.5	25 U	50 U	
Α	5106-000921-053	B-69	9/21/00	10	25 U	50 U	
A	5106-000921-055	B-69	9/21/00	20	25 U	50 U	
Α	5106-000921-056	B-70	9/21/00	2.5	25 U	50 U	
Α	5106-000921-057	B-70	9/21/00	10	25 U	50 U	
Α	5106-000921-058	B-70	9/21/00	20	25 U	50 U	•
A	5106-000921-059	B-70	9/21/00	26.5	612	523	
Ā	5106-000921-060	B-71	9/21/00	2.5	25 U	50 U	
Α	5106-000921-061	B-71	9/21/00	10	25 U	. 50 U	,
Α	5106-000921-062	B-71	9/21/00	20	25_U	50 U	
A	5106-000921-063	B-72	9/21/00	2.5	25 U	84	
Α	5106-000921-064	B-72	9/21/00	12.5	25 U	126	
A	5106-000921-065	B-72	9/21/00	20	25 U	50 U	
^_	—5106-000921-066—	B-72 	9/21/00	—2 .5 —	25 U	50 U 50-U	
	5106-00 0921-067		9/21/00		25 U	50-U	
Â	5106-000921-068	B-73	9/21/00	12.5	25 U	50 U	
Â	5106-000925-079	B-73 B-74	9/25/00	2.5			
- Â	5106-000925-080				25 U	50 U	
	5106-000925-081	B-74	9/25/00	12.5	25 U	50 U	_
A	5106-000925-082	B-74	9/25/00	18	25 U	50 U	
		B-78	9/25/00	20	25 U	50 U	•
A	5106-000921-075	B-81	9/22/00	2.5	25 U	50 U	
A	5106-000921-076	B-81	9/22/00	10	68.2	90.5	
A	5106-000921-077	B-81	9/22/00	20	864	2020	
A	5106-000921-070	B-82	9/22/00	2.5	250 U	673	
A	5106-000921-071	B-82	9/22/00	10	250 U	1570	
A	5106-000921-072	B-82	9/22/00	20	787	846	
Α	5106-000921-074	B-82	9/22/00	26	25 U	50 U	
A	5106-000920-038	B-84	9/20/00	67.5	107	191	
_A	5106-000921-078	B-86	9/22/00	10	25 U	50 U	
_A	—5106-001030-08 1 —	—В-94——	10/30/00 -	2:5	506 U	1300	
Α	5106-001030-082	B-94	10/30/00	10	24.4 U	61 U	
Α	5106-001030-083	B-94	10/30/00	20	28.6 U	71.4 U	
-A	5106-001030-084	B-94	10/30/00	25.5	28.6 U	71.4 U	
A .	5106-001106-095	B-97	11/6/00	- 3	24.7 U	61.7 U	
Α	5106-001106-096	B-97	11/6/00	5	26 U	64.9 U	

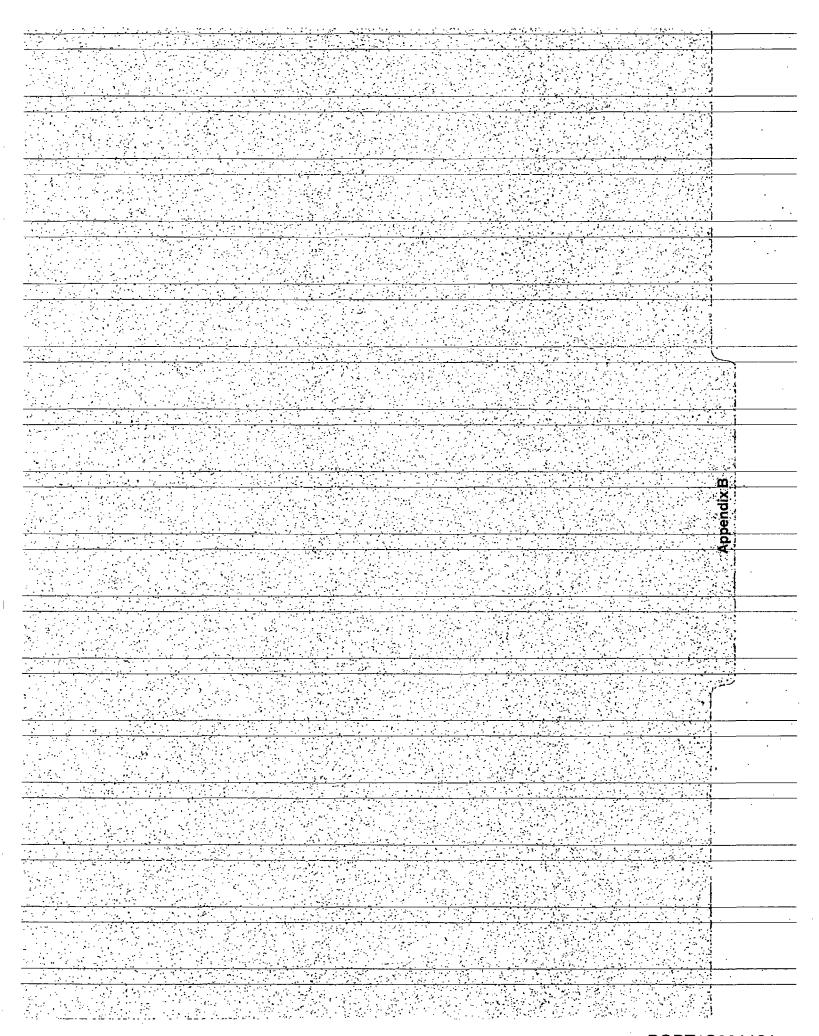
Area	Sample	Station	Sample	Depth	NW Method	TPH-D in mg/kg	
	Number		Date	in feet		Oil	
_A	5106-001106-097	B-97	11/6/00	10	26.7 U	66.7 U	
Α	5106-001106-098	B-97	11/6/00	20	197	266	
Α	5106-001106-099	B-97	11/6/00	26	28.6 U	71.4 U	
Α	5106-001106-090	B-99	11/6/00	3	24.4 U	61 U	
_A:	5106-001106-092	_B-99	11/6/00	10	27.4 U	68.5 U	
A	5106-001106-093	B-99	11/6/00	- 20-	28.2-U-	70.4 U	
Α	5106-001106-094	B-99	11/6/00	26	24.4 U	61 U	
В	5106-001024-008	B-106	10/24/00	7	25 U	174	
-B	5106 -001024 - 009	B-106	10/24/00	16.5	250		
В	5106-001024-011	B-106	10/24/00		25 U	50 U	·
В	4876-000229-003	B-15	2/29/00	2	25.UR	50°UR	
В	4876-000302-028	B-16	3/2/00	1.5	25 UR	* * * * * * * * * * * * * * * * * * *	
В	4876-000301-014	B-21	3/1/00	1.5	25 UR	50 UR A	
В	4876-000301-011	B-22	3/1/00	1.5	25 R	50 R	
В	4876-000229-008	B-29	2/29/00	4	725 UE	50 UR	
В	4876-000301-013	B-33	3/1/00	1.5	25 25 UK	50 UR 5	
В	4876-000301-012	B-34	3/1/00	1.5	25 UR		
В	4876-000301-016	B-37	3/1/00	10.5		300K	
В	T-1 B-4 0-2	B-4	3/26/98	0-2	25 U	50 U	
В	T-1 B-5 0-2	B-5	3/26/98	0-2	500 U	6030	
В	5106-000919-019	B-58	9/19/00	8	25 U	50_U_	
В	5106-000919-020	B-59	9/19/00	4	25 U	112	
В	T-1 B-6 (0-2)	B-6	3/26/98	0-2	25 U	50 U	•
_B	5106-000919-017	—B-60——	9/19/00	_4_	25 U	50 U	
_В	5106-000919-016	—B-61—	9/19/00 -	4	25 U	50 U	
В	5106-000919-021	B-62	9/19/00	4	25 U	50 U	
В	B-63-Duplicate	B-63	9/20/00	10.5	1170	3210	
В	5106-000920-022	B-63/63a	9/20/00	6	500 U	1190	
B	5106-000920-023	B-63/63a	9/20/00	10.5	3440	10000	
В	5106-000920-024	B-63/63a	9/20/00	16	250 U	2180	
В	5106-001024-054	B-63/63a	10/25/00	19	98.3	286	
В	5106-001024-055	B-63/63a	10/25/00	24	25 U	50 U	
В	5106-000919-005	B-64/64a	9/19/00	10.5	25 U	50 U	
В	5106-001024-037	B-64/64a	10/25/00	2.5	25 U	50 U	
В	5106-001024-040	B-64/64a	10/25/00	16.5	109	251	
В	5106-001024-041	B-64/64a	10/25/00	19	39	105	
В	5106-000919-006	B-65/65a	9/19/00	10.5	250 U	769	
В	5106-000979-000	B-65/65a	9/25/00	16.5	500 U	9070	
.B	5106-000925-084	B-65/65a	9/25/00 9/25/00	19.5	25_U	9070 50 U	
-B	5106-001024-043	— В-05/65а — —В-65/65а—	<i>9125/00-</i> 10/25/00-		250-U	30-U 1170	
-Б В	5106-001024-046	—в-65/65а— В-65/65а	10/25/00 10/25/00	— 2.5 — · 12	2500 U	20700	
В	5106-001024-046	B-66/66a	9/19/00				
B	-5106-000919-007	B-66/66a	9/19/00 9/ 19/00	10.5 - 18	1090	2380	
-B	5106-000919-008 5106-000919-009	B-66/66a	9/19/00		-3830	6320	
В	5106-001024-047	B-66/66a	10/25/00	19 2.5	250 U	217 1650	

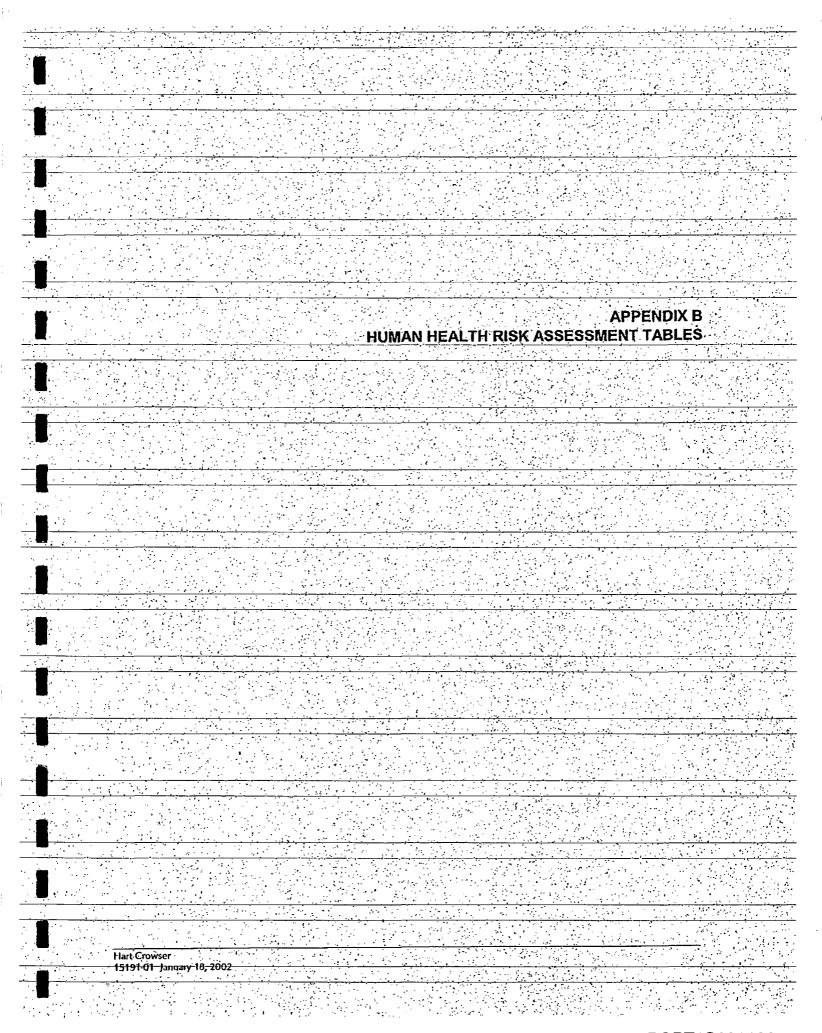
Area	Sample	Station	Sample	Depth	NW Method	TPH-D in mg/kg	
	Number		Date	in feet	Diesel	Oil	
В	5106-000919-015	B-67/67a	9/19/00	13.5	29	62.7	
В	5106-001030-085	B-67/67a	10/30/00	5.5	20.8 U	52.1 U	
В	5106-001030-086	B-67/67a	10/30/00	13.5	28.6 U	124	
В	5106-001030-087	B-67/67a	10/30/00	19.5	24.1 U	60.2 U	
B	5106-001030-089	B-67/67a	10/30/00	23.5	46.4	203	
B	<u> 5106-000919-010</u>	—B-83	9/19/00	4	25 U	50-U	
В	5106-000919-014	B-83	9/19/00	12	25 U	50 U	
В	5106-000920-025	B-85	9/20/00	12.5	1060	3000	
B	5106 -000 920-026	B=85	9/20/00-	19	1310	2640	
В	5106- 00092 0-028	B-85	9/20/00	24	25 U	50 U	
В	5106-001024-032	B-87	10/25/00	2.5	25 U	1860	
В	5106-001024-034	B-87	10/25/00	10.5	25 U	50 U	
<u></u> В	5106-001024-035	B-87	10/25/00	19	25 U	98.7	
-В	5106-001024-036	B-87	10/25/00	24	25 U	144	
В	5106-001024-027	B-88	10/25/00	6.5	25 U	148	
В	5106-001024-029	B-88	10/25/00	16.5	49	164	
В	5106-001024-030	B-88	10/25/00	19	25 U	50 U	
В	5106-001024-031	B-88	10/25/00	24	46	50 U	
В	5106-001024-021	B-89	10/24/00	2.5	25 U	50 U	
В	5106-001024-023	B-89	10/24/00		25 U	50_U	
B	5106-001024-024	B-89	10/24/00	19	45.7	82,5	
В	5106-001024-025	B-89	10/24/00	24	25 U	50 U	
В	5106-001024-018	B-90	10/24/00	16.5	25 U	50 U	
В	5106-001024-019	B-90	10/24/00	19	-324	1150	
В —	—5106-00 1024-020 —	-B-90	10/24/00	24-	-35.9	61.8	
В	B-90-Duplicate	B-90	10/24/00	16.5	25 U	50 U	
В	5106-001024-001	B-91	10/24/00	2.5	25 U	50 U	
-B	5106-001024-005	B-91	10/24/00		250 U	1150	
_B	5106-001024-006	B-91	10/24/00	19	25 U	50 U	
В	B-91-Duplicate	B-91	10/24/00	2.5	25 U	50 U	
В	5106-001024-013	B-92	10/24/00	10	169	310	
В	5106-001024-015	B-92	10/24/00	17.5	250 U	1970	·
						sment/Appendicies/App-A	

J = Estimated Concentration.
 R = Rejected Data (see Appendix F).
 Shading indicates Rejected Data.

Sample ID Station	5108-011001-108 MW-1	5106-011001-109 MW-1 (Dup)	5106-011001-107 MW-2	5106-010928-103 MW-3	5106-010928-104 MW-4
Sampling Date	10/01/2001	10/01/2001	10/01/2001	9/28/2001	9/28/2001
Depth in Feet	17 - 32	17 - 32	17 - 32	17 - 32	17 - 32
Total Suspended Solids	36	35	55	720	130
Total Metals In μg/L	•				
Arsenic	2.01	1.06	12.8	14	6.45
Cadmium	1 U	1 U			1 U
Chromium	3.25	2.65			5.12
Copper	4.74	3.88	2 U	40.2	4.48
Lead	1.16	1 U	1 U	36.2	2.49
Mercury	0.2 U	0.2 U			0.2 U
Nickel	5.25	4.49			3.86
Silver	1-⊍	———1-U	<u>:</u>		1-U
Zinc	10.6	8.43			9.06
Dissolved Metals in µg/L	**		•	•	
Arsenic	1 U	1 U	14.5	11	6.51
Copper	2.29	2.03	2_U	2 U	2 U
Lead	1.37	1 U	1 U	1_U	1.0
PAHs in μg/L					W. W. C. C. C. C. C. C. C. C. C. C. C. C. C.
Benzo(a)anthracene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U
Benzo(a)pyrene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U
Benzo(b)fluoranthene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 ป
Benzo(k)fluoranthene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U
Chrysene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U
Dibenz(ah)anthracene	0.2 U	0.2 U	0.2 U	0.2 U	0.2 U
Indeno(1,2,3-cd)pyrene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U
Acenaphthene	0.1 U	0.1 U	0.121	0.192	0.72
Acenaphthylene	0.1 U	0.1 U	0.121 0.1 U	0.132 0.1 U	0.1 U
Anthracene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U
Benzo(ghi)perylene	0.1 U	0.1 U	0.1 U	0.1 U	0.1 U
Fluoranthene	0.1 U	0.1 U	0.10	0.1 U	0.1 U
Fluorene	0.1 U	0.1 U	0.119 0.1 U	0.1 U	0.1 U
Naphthalene	0.1 U	0.1 U	0.1 U	0.1 ป 0.1 ป	0.1 0
Phenanthrene	0.1-U	0.1 U 	1.25	0.1 U 0 .138	•
Pyrene Pyrene	0.1 U		0.564		
Total PAHs	0.2 U				
TPH in µg/L	U.Z U	0.2 U	2.054	0.33	1.71
• •	416.2	220 0			050 11
Diesel		338.2			250 U
Oil Malatiles in well	500 U	500 U			500 U
Volatiles in µg/L Tetrachloroethene	2.76	3.29			1.0 U
•					

Table A-3 - Analytical	Results for W	later Samples			Sheet 2 of 2
Sample ID	5108-010928-102	5106-010928-105	5106-011001-106		
Station	MW-5	MW-6	MW-7.	·	
Sampling Date	9/28/2001	9/28/2001	10/01/2001		
Depth in Feet	19 - 34	17 - 32	17 - 32		
Total Suspended Solids	108	50	20 U	•	
Total Metals in µg/L					
Arsenic	12.1	2.72	1.38	·	
Cadmium					
Chromium					
Copper	2.95	2.51	2 U		
Lead	1.48	1 U	4.47		· · · · · · · · · · · · · · · · · · ·
Mercury					
Nickel					
Silver		_			
Zinc	· ·		11.6	<u> </u>	
Dissolved Metals in µg/L					
Arsenic	11.3	3.65	1 U		
Copper	2_U	2 U	2U		
Lead	1 U	. 1 U	1.0		
PAHs in µg/L		•			
Benzo(a)anthracene	0.1 U	.0.1 U	0.1 U		
Benzo(a)pyrene		0.1 U	0.1 U		
Benzo(b)fluoranthene	0.1 U	0.1 U	0.1 U		
Benzo(k)fluoranthene	0.1 U	0.1 U	0.1 U		
Chrysene	0.1 U	0.1 U	0.1 U		
Dibenz(ah)anthracene	0.2 U	0.2 U	0.2 U		
Indeno(1,2,3-cd)pyrene	0.1 U	0.1 U	0.1 U		
Acenaphthene	0.448	0.1 U	0.1 U		
Acenaphthylene	0.1 U	0.1 U	0.1 U		
Anthracene	0.1 U	0.1 U	0.1 U		
Benzo(ghi)perylene	0:1 ⁻ U	0.1 U	0:1 U	-	• • • • • • • • • • • • • • • • • • • •
Fluoranthene	0.1 U	0.1 U	0:1·U		
Fluorene	0.1 ป	0.1 U	0.1 U		•
Naphthalene	0.1 U	0.1 U	0.1 U		
Phenanthrene	1.16	0:1-U	0.153	•	
	0.172				
Total PAHs	1.78	0.2 U	0.306		
TPH in μg/L				•	
Diesel		<u> </u>			
Qil					
Volatiles in µg/L					
Tetrachloroethene	•				
	Notes:	F:VDAT	AU006/Fort of Portland\15191-01 T-	1 Risk Assessment/Appendictes/App-A	
•		cted at ar above t	he method reporting li	mits	
		ed Concentration.	moasou reponing ii	inio.	
		Data (see Apper	ndix F)		<u>-</u>
	o. n - nejeuleu	Daia loce whitei	iuin i ji		·
				,	•
				<u> </u>	<u> </u>
					
			•	•	
		•			





Sheet 2 of 4

Table B-1 - Area A Risk Calculations

Table B-2 - Area A Risk Calculations Soil Ingestion, Commercial Worker Marine Terminal 1 South Risk Assessment Portland, Oregon

Compounds of Potential	Soll EPC	in mg/kg	II	Intake in g-day	Hazard	Quotient	6 1 1	Intake in g-day	Cance	r Risk
Concern	RME	СТ	RME	ст	RME	ст	RME	СТ	RME	ст
PAHs										
Benzo(a)anthracene	2.0E+00	6.7E-01	2.0E-06	3.3E-07			7.0∉-07	2.8E-08	5.1E-07	2.1E-08
Benzo(a)pyrene	1,8E+00	5.5E-01	1.8E-06	2.7E-07			6.3∉-07	2.3E-08	4.6E-06	1.7E-07
Benzo(b)fluoranthene	1.4E+00	4.0E-01	1.4E-06	2.0E-07			4.9 Ė- 07	1.7E-08	3.6E-07	1.2E-08
Dibenz(a,h)anthracene	1 6E-01	9.0E-02	1.6E-07	4.4E-08		-	5.6E-08	3.8E-09	4.1E-07	2.8E-08
ndend(1,2,3-cd)pyrene	7 4E-01	2.8E-01	7.2E-07	1.4E-07	-	-	2.6ᡛ-07	1.2E-08	1.9E-07	8.6E-09
Arsenic	8.4E+00	3.3E+00	8.2E-06	1.6E-06	2.7E-02	5.4E-03	2.91 -06	1.4E-07	4.4E-06	2.1E-07
			TOTAL HAZ	ARD INDEX	3.E-02	5.E-03	TOTAL CAL	NCER RISK	1.E-05	4.E-07

Notes:

RME = Reasonable Maximum Exposure.
CT = Central Tendency.
EPC = Exposure Point Concentration.

Hazard Intake in

mg/kg-day

Hazard Quotient

Sheet 2 of 4

Cancer Risk

Cancer Intake in

mg/kg-day

Table B-2 - Area A Risk Calculations

Portland, Oregon

Compounds of Potential

Dermal Contact with Soil, Commercial Worker Marine Terminal 1 South Risk Assessment

Soil EPC in mg/kg

Table B-2 - Area A Risk Calculations Fugitive Dust Inhalation (Outdoor Air), Commercial Worker
Marine Terminal 1 South Risk Assessment Portland, Oregon

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		Outdoor Air EP mg/m³		Taken a meane in		Hazard Quotient		Cancer I mg/k	1 1	Cancer Risk	
Compounds of Potential Concern	PEF in m³/kg	RME	ст	RME	ст	RME	СТ	RME	ст	RME	СТ
'AHs									1 1		
Benzo(a)anthracene	1.325+09	1.5E-09	5.1E-10	2.3E-10	7.5E-11			8.0E-11	6.5E-12	2.5E-11	2 0E-12
Benzo(a)pyrene	1.32€+09	1.4E-09	4.2E-10	2.0E-10	6.2E-11	_	-	7.2E-11	5.3E-12	2.2E-10	1,6E-11
Benzo(b)fluoranthene	1.32E+09	1.1E-09	3.0E-10	1.6E-10	4.5E-11			5.6E-11	3.9E-12	1.7E-11	1 2E-12
Dibenz(a,h)anthracene	1.325+09	1.2E 10	6.8E-11	1.8E-11	1.0E-11		\ _	6.4E-12	8.7E-13	2.0E-11	27E-12
ndeno(1,2,3-cd)pyrene Vietals	1.325+09	5.6E-10	2. E-10	8.3E-11	3.2E-11			3.0E-11	2.7E-12	9.2E-12	8.4E-13
Arsenic	1.32E+09	6.4E-09	2.5E-09	9.5E-10	3.7E-10	_		3.4E-10	3.2E-11	5.1E-09	4.8E-10
				TOTAL HAZ	ARD INDEX	0.E+00	0.E+00	TOTAL CAN	CER RISK	5.E-09	5.E-10

Outdoor Air EPC = Soil EPC (See Table 4)/PEF.
PEF = Particulate Emission Factor
RME = Reasonable Maximum Exposure.
CT = Central Tendency.
EPC = Exposure Point Concentration.

F: DATA Vobs Port of Portand 16191-01 T-1 Risk Assessment Appendices Appendix B Tables B-2

Hazard Intake in

Sheet 1 of 4

Cancer Intake in

Table B-3 - Area A Risk Calculations

Marine Terminal 1 South Risk Assessment

Soil Ingestion, Excavation Worker

Portland, Oregon

Table B-3 - A	rea A Risi	Calculati	ons
Dermai Cont	tact with \$	oil, Excav	ation Worker
Marine Term	inal 1 Sou	th Risk As	sessment
Portland, Or	egon		

		Cance	r Ri	sk	
		RME		СТ	
	,				
0	7.	2E-10	8	.5E-11	H
Λ I	-	0- 00	11 -7	CT 40	и

Sheet 2 of 4

Compounds of Potential		Soil EPC	in mg/kg	1 1 1	Intake in g-day	Hazard (Quotient	Cancer In mg/kg-		Cance	Risk
Concern	ABS	RME	СТ	RME	СТ	RME	CT	RME	СТ	RME	СТ
PAHs											
Benzo(a)anthracene	0.13	3.7E+01	3.7E-01	6.9E-08	1.6E-08			9.9E-10	1.2E-10	7.2E-10	8.5E-11
Benzo(a)pyrene	0,13	3.7E-01	3.3E-01	6.9E-08	1.5E-08		·	9.9E-10	1.0E-10	7.2E-09	7.6E-10
Benzo(b)fluoranthene	0.13	3.4E-01	2.7E-01	6.4E-08	1.2E-08	-	1	9.1E-10	8.5E-11	6.7E-10	6.2E-11
Dibenz(a,h)anthracene	0 13	7.0E-02	6.0E-02	1.3E-08	2.6E-09	4		1.9E-10	1.9E-11	1.4E-09	1.4E-10
Indeno(1,2,3-cd)pyrene Metals	0.13	2.0E-01	1.8≣-01	3.8E-08	7.9E-09		-	5.4E-10	5.7E-11	3.9E-10	4.1E-11
Arsenic	0.03	6.0E+00	3.4E+00	2.6E-07	3.4E-08	8.7E-04	1.1E-04	3.7E-09	2.5E-10	5.6E-09	3.7E-10
				TOTAL HAZ	ARD INDEX	9.E-04	1.E-04	TOTAL CANC	ER RISK	2.E-08	1.E-09

Notes:

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ABS = Dermal Absorption Factor (EPA, 1998).
RME = Reasonable Maximum Exposure.
CT = Central Tendency,
EPC = Exposure Point Concentration.

Sheet 3 of #

Table B-3 - Area A Risk Calculations

		Outdoor A mg	_	Hazard I mg/kg	1	Hazard Q	uotient	Cancer Int		Cancer	Risk
Concern Concern	PEF in m³/kg	RME	ст	RME	ст	RME	ст	RME	ст	RME	ст
AHS											
Benzo(a)anthracene	1.32E+09	2.8E-10	2.8E-10	1.5E-12	1.5E-12	\ <u>-</u> \	1 1	2.1E-14	1.1E-14	6.6E-15	3 3E-15
lenzo(a)pyrene	1.325+09	2.8E-10	2.5E-10	1.5E-12	1.3E-12	-		2.1E-14	9.6E-15	6.6E-14	3.0E-14
enzo(b)fluoranthene	1.32E+09	2.6E-10	2.0E-10	1.4E-12	1.1E-12	- -		2.0E-14	7.8E-15	6.1E-15	2:4E-15
benz(a,h)anthracene	1.32日+09	5.8E-11	4.5E-11	2.8E-13	2.4E-13			4.1E-15	1.7€-15	1.3E-14	5.4E-15
ndeno(1,2,3-cd)pyrene Metals	1.32E+09	1.5E-10	1.4E-10	8.1E-13	7.3E-13	-	1 +	1.2E-14	5.2E-15	3.6E-15	1.6E-15
vrsenic	1.32E+09	4.5E-09	2.6E-09	2.4E-11	1.4E-11		4	3.5E-13	9.9∉-14	5.2E-12	1.5E-12
				TOTAL HAZ	ARD INDEX	0.E+00	0.E+00	TOTAL CANCE	R RISK	5.E-12	2.E-12

Outdoor Air EPC = Soll EPC (See Table 4)/PEF.
PEF = Particulate Emission Factor
RME = Reasonable Maximum Exposure.
CT = Central Tendency.
EPC = Exposure Point Concentration.

F:DATALlobs Port of Portand 15191-01 T-1 Risk Assessment Appendictes (Appendix B Tables B-3

Table B-4 - Area B Risk Calculations Soil Ingestion, Resident Marine Terminal 1 South Risk Assessment Portland, Oregon

	Soll EPC	in mg/kg	11	ntake in g-day	Hazard	Quotient	[[Intake in g-day	Cance	r Risk
Compounds of Potential Concern	RME	ст	RME	СТ	RME	CT	RME	ст	RME	СТ
PAHs	,			-						
Benzo(a)anthracene	1.5E-01	6.0E-02	3.8E-06	4.4E-08			4.0E-07	4.0E-09	2.9E-07	2 9E-09
Benzo(a)pyrene	1.9E-01	7.0E-02	4.9E-06	5.1E-08	_	4	5.1E-07	4.6E-09	3.7E-06	3.4E-08
Benzo(b)fluoranthene	1.3E-01	6.0E-02	3.3E-06	4.4E-08		\ \\\	3.5E-07	4.0E-09	2.5E-07	2.9E-09
Indeno(1,2,3-cd)pyrene	1.2E-01	5.0E-02	3.1E-06	3.7E-08	-	+ 1	3.2E-07	3.3E-09	2.3E-07	2.45-09
Arsenic	3.1E+00	2.9E+00	7.9E-05	2.1E-06	2.6E-01	7.1Ε-03	8.3E-06	1.9E-07	1.2E-05	2.9E-07
	1 1		TOTAL HAZ	ARD INDEX	3.E-01	7.E-03	TOTAL CAN	CER RISK	2.E-05	3.E-07

Notes:

RME = Reasonable Maximum Exposure.
CT = Central Tendency.
EPC = Exposure Point Concentration.

Table B-4 - Area B	Risk Calcul	lations	
Fugitive Dust Inhal	ation (Outd	loor Air), Res	ident
Marine Terminal 1	South Risk	Assessment	
Portland, Oregon			

Sh	eet	3	of	3
----	-----	---	----	---

		1	Air EPC in _I /m ³	Hazard II mg/kg		Hazard Q	uctient	Cancer III		Cance	r Risk
Compounds of Potential Concern	PEF in m³/kg	RME	ст	RME	ст	RME	СТ	RME	ст	RME	СТ
РАН Б	.										
Benzo(a)anthracene	1.32E+09	1.1E-10	4.5E-11	6.0E-11	2.4E-11	-		2.9E-11	2.5E-12	8.9E-12	7.7E-13
Benzo(a)pyrene	1.825+09	1.4E-10	5. 3E- 11	7.6E-11	2.8E-11		1 1	3.6E-11	2.9E-12	1.1E-10	8.9E-12
Benzo(b)fluoranthene	1.82E+09	9.8E-11	4.5E-11	5.2E-11	2.4E-11		.	2.5E-11	2.5E-12	7.7E-12	7.7E-13
ndeno(1,2,3-cd)pyrene	1.325+09	9.1E-11	3.8E-11	4.8E-11	2.0E-11		1	2.3E-11	2.1E-12	7.1E-12	6.4E-13
Vietals					j						
Arsenic	1.82E+09	2.3E-09	2.2E-09	1.2E-09	1.2E-09		↓ ↓	5.9E-10	1.2E-10	8.9E-09	1.8E-09
		- 1		TOTAL HAZ	ARD INDEX	0,E+00	0.E+00	TOTAL CAN	CER RISK	9.E-09	2.E-09

Outdoor Air EPC = Soil EPC (See Table 4)/PEF.
PEF = Particulate Emission Factor.

RME = Reasonable Maximum Exposure.

CT = Central Tendency

EPC - Exposure Point Concentration.

F: DATA Jobs Port of Pontand 15191-01 T-1 Risk Assessment Appendictes Appendix 8 Tables 18-4

Table B-5 - Area B Risk Calculations Fugitive Dust Inhalation (Outdoor Air), Commercial Worker Marine Terminal 1 South Risk Assessment Portland, Oregon

Sheet 3 of B

			Air EPC in s/m³		d Intake in /kg-day	Hazard Quotient	Cancer Intaké in mg/kg-day	Cancer Ris	k
Compounds of Potential Concern	PEF in m³/kg	RME	ст	RME	ст	RME CT	RME CT	RME	ст
AHs									
Benzo(a)anthracene	1.325+09	1.1E-10	4.5E-11	1.7E-11	6.8E-12		6.0E-12 5.8E-13	1.95-12 1.9	8E-13
Benzo(a)pyrene	1.325+09	1.4E-10	5.3E-11	2.1E-11	7.9E-12		7.6E-12 6.8E-13	2.4E-11 2.	1E-12
Benzo(b)fluoranthene	1.32E+09	9.BE-11	4.5E-11	1.5E-11	6.8E-12	- +	5.2E-12 5.8E-13	1.6E-12 1.	8E-13
ndeno(1,2,3-cd)pyrene	1.32E+09	9.1E-11	3.8E-11	1.4E-11	5.6E-12	. +	4.8E-12 4.8E-13	1.55-12	5E-13
rsenic	1.325+09	2.3E-09	2.2E-09	3.5E-10	3.3E-10		1.2E-10 2.8E-11	1.9E-09 4.:	2E-10
				TOTAL H	AZARD INDEX	0.E+00 0.E+0	O TOTAL CANCER RISK	2.E-09 4	.E-10

Notes:

Outdoor Air EPC = Soil EPC (See Table 4)/PEF.
PEF = Particulate Emission Factor.
RME = Reasonable Maximum Exposure.

CT = Central Tendency.
EPC = Exposure Point Concentration.

Table B								
Derma	Conta	ıct w	ith So	il, Exc	av	ation V	Vo	rker
Marine	Termi	nal 1	Sout	h Risk	A۶	sessm	er	it
Portlar	id, Ore	gon						

Compounds of Potential		Soli EPC in mg/kg		Hazard Intake in mg/kg-day		Hazard Quotient		Cancer Intake in mg/kg-day		Cancer Risk	
	ABS	RME	ст	RME	ст	RME	ст	RME	ст	RME	ст
PAHs		.									
Benzo(a)anthracene	0.13	1.5E+00	4.0E-01	2.8E-07	1.8E-08	-	-	4.0E-09	1.3E-10	2.9E-09	9.2E-11
Benzo(a)pyrene	0.13	2.1E+00	4.7E-01	3.9E-07	2.1E-08			5.6E-09	1.5E-10	4.1E-08	1.1E-09
Benzo(b)fluoranthene	0 13	1.4E+00	3.7E-01	2.6E-07	1.6E-08	-	-	3.8E-09	1.2E-10	2,7E-09	8.5E-11
Dibenz(a,h)anthracene	0,13	2.3E-01	1.1E-01	4.3E-08	4.8E-09			6.2E-10	3.5E-11	4.5E-09	2.5E-10
Indeno(1,2,3-cd)pyrene Metals	0.13	9.0E-01	2.9E-01	1.7E-07	1.3E-08		-	2.4E-09	9.1E-11	1.8E-09	6.6E-11
Arsenic	0.03	3.3E+00	2.9E+00	1.4E-07	2.9E-08	4.8E-04	9.8E-05	2.0E-09	2.1E-10	3.1E-09	3.2E-10
				TOTAL HAZ	ARD INDEX	5.E-04	1.E-04	TOTAL CAN	CER RISK	6.E-08	2.E-09

Sheet 2 of 3

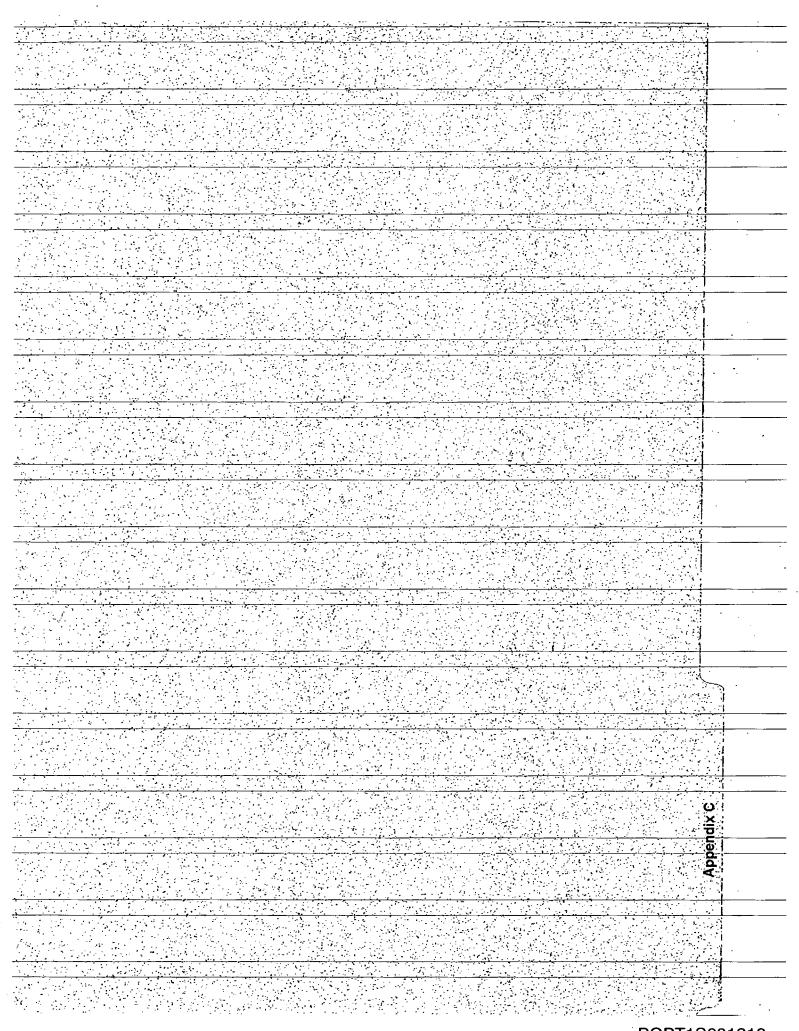
ABS = Dermal Absorption Factor (EPA, 1998).
RME = Reasonable Maximum Exposure.
CT = Central Tendency.
EPC = Exposure Point Concentration.

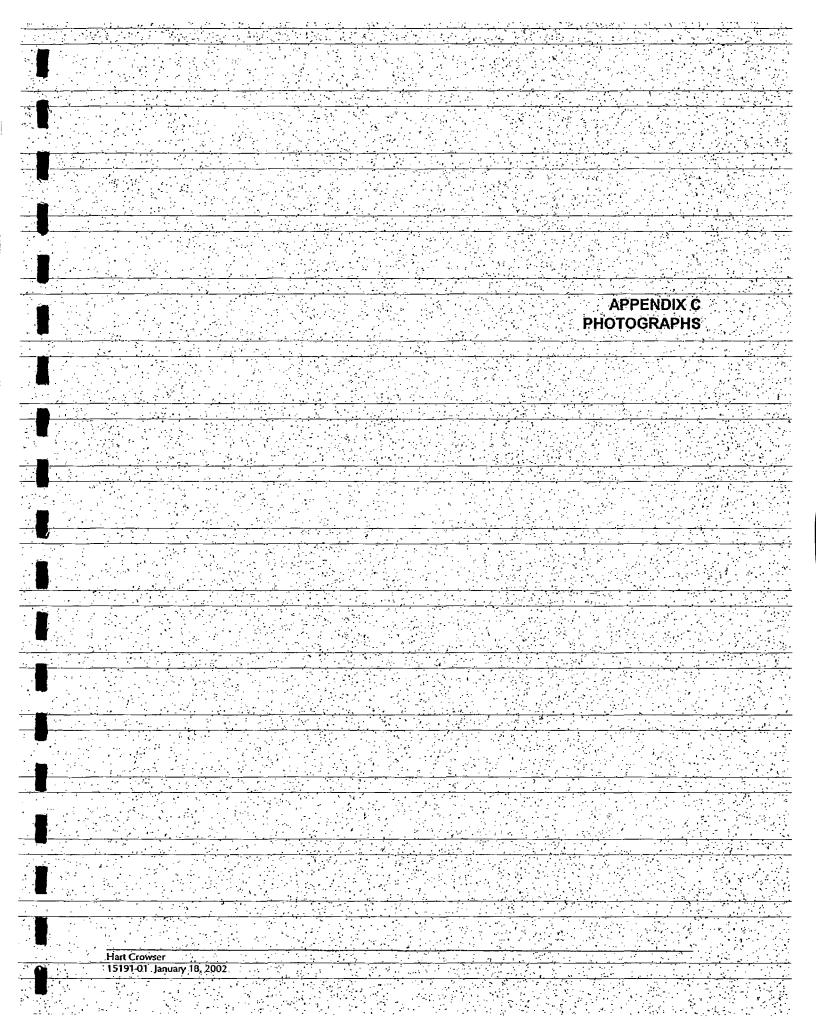
Table B-6 - Area E	Risk Calcu	lations				
Fugitive Dust Inha	lation (Outo	ioor Air),	Ėχ	avatio	n١	Worker
Marine Terminal 1	South Risk	Assessm	en	t		
Portland, Oregon						

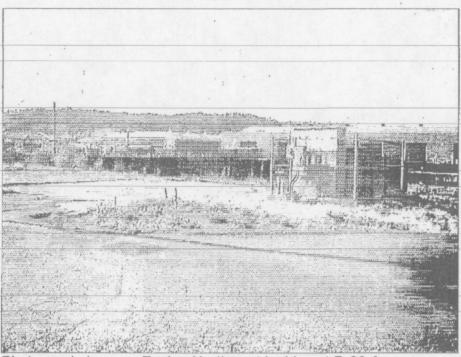
		1	Air EPC in	Hazard Intake In mg/kg-day		Hazard Quotient		Cancer Intake in mg/kg-day		Cancer Risk	
ompounds of Potential Concern	PEF in m³/kg	RME	ст	RME	СТ	RME	ст	RME	ст	RME	ст
AHs											
enzo(a)anthracene	13E+09	1.1E-09	3.0E-10	6.1E-12	1.6E-12	_	1	8.7E-14	1.2E-14	2.7E-14	3.6E-15
enzo(a)pyrene	1.3E+09	1.6E 09	3.6E-10	8.5E-12	1.9E-12			1.2E-13	1.4E-14	3.8E-13	4.2E-14
enzo(b)fluoranthene	13E+09	1.1E-09	2.8E-10	5.7E-12	1.5E-12	· -	1	8.1E-14	1:1E-14	2.5E-14	3.3E-15
ibenz(a,h)anthracene	1,3E+09	1.7E-10	8.3E-11	9.3E-13	4.5E-13	-	+	1.3E-14	3.2E-15	4.1E-14	9.9E-15
deno(1,2,3-cd)pyrene	1.3E+09	6.BE-10	2.2E-10	3.7E-12	1.2E-12		4	5.2E-14	8.4E-15	1.6E-14	2.6E-15
etals											
rsenic	1.3E+09	2.5E-09	2.2E-09	1.3E-11	1.2E-11	-		1.9E-13	8.4E-14	2.9E-12	1.3E-12
				TOTAL HAZ	ARD INDEX	0.E+00	0.E+00	TOTAL CAN	CER RISK	3.E-12	1.E-12

Sheet 3 of 3

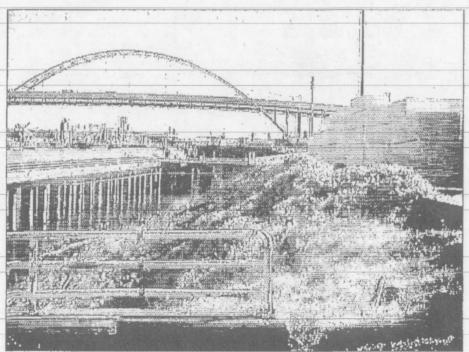
Outdoor Air EPC = Soil EPC (See Table 4)/PEF.
PEF = Particulate Emission Factor
RME = Reasonable Maximum Exposure.
CT = Central Tendency.
EPC = Exposure Point Concentration.



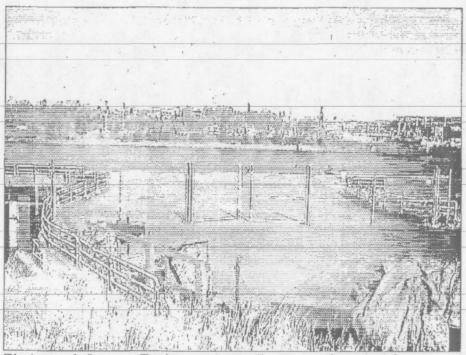




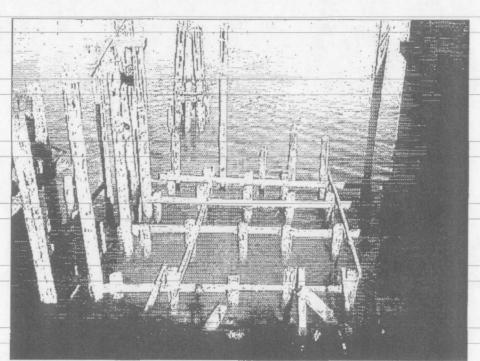
Photograph 1- Facing Northwest looking at B-20 Area.



Photograph 2 - Facing Southeast looking at bank area near Concrete Pier.



Photograph 3 - Facing east standing at south end of Concrete Pier.



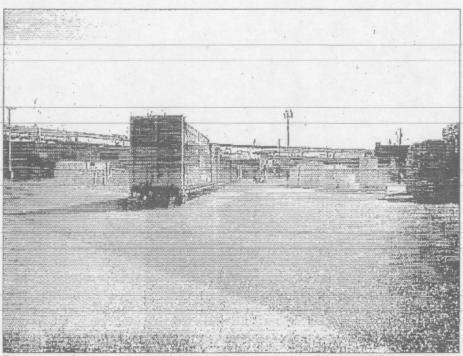
Photograph 4 - Facing east standing south of Concrete Pier.



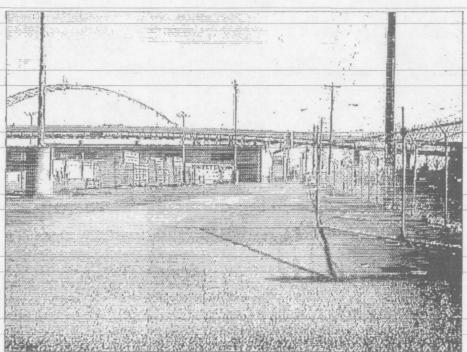
Photograph 5 - Facing west viewing north end of Terminal 1.



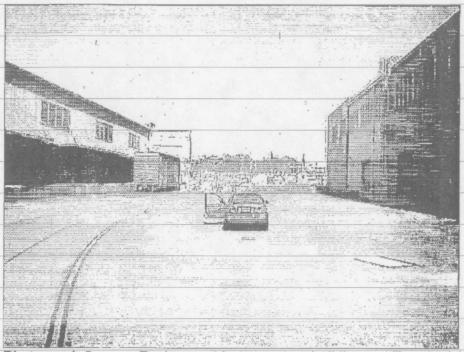
Photograph 6 - Facing southwest viewing north end of Terminal 1.



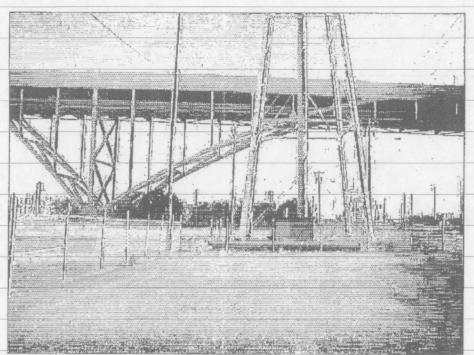
Photograph 7 - Facing south viewing north end of Terminal 1.



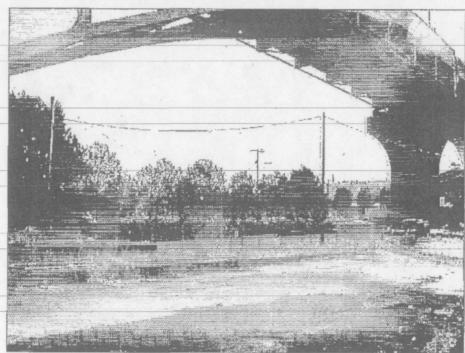
Photograph 8 - Facing south viewing north end of Terminal 1.



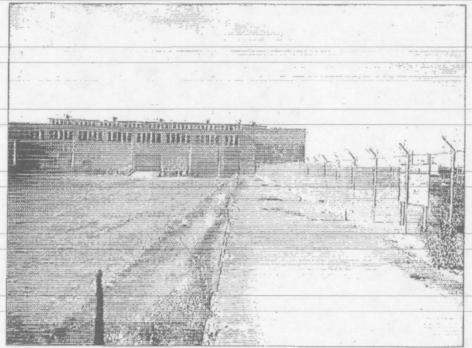
Photograph 9 - Facing east between two buildings at Terminal 1.



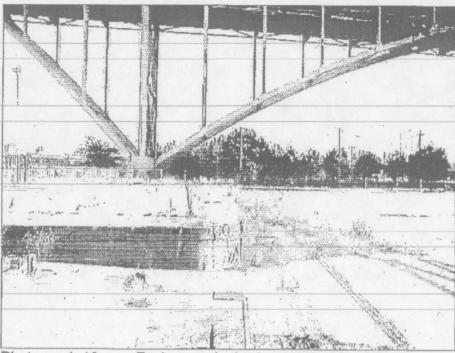
Photograph 10 - Facing south viewing south end of Terminal 1.



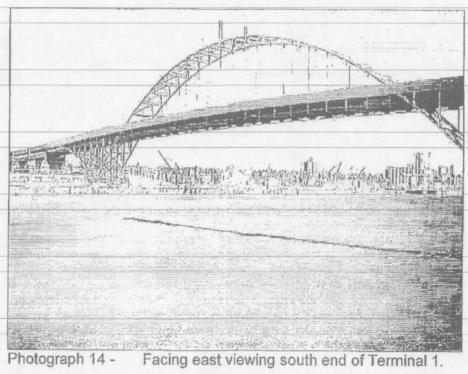
Photograph 11 - Facing south viewing adjacent property at south end of Terminal 1.



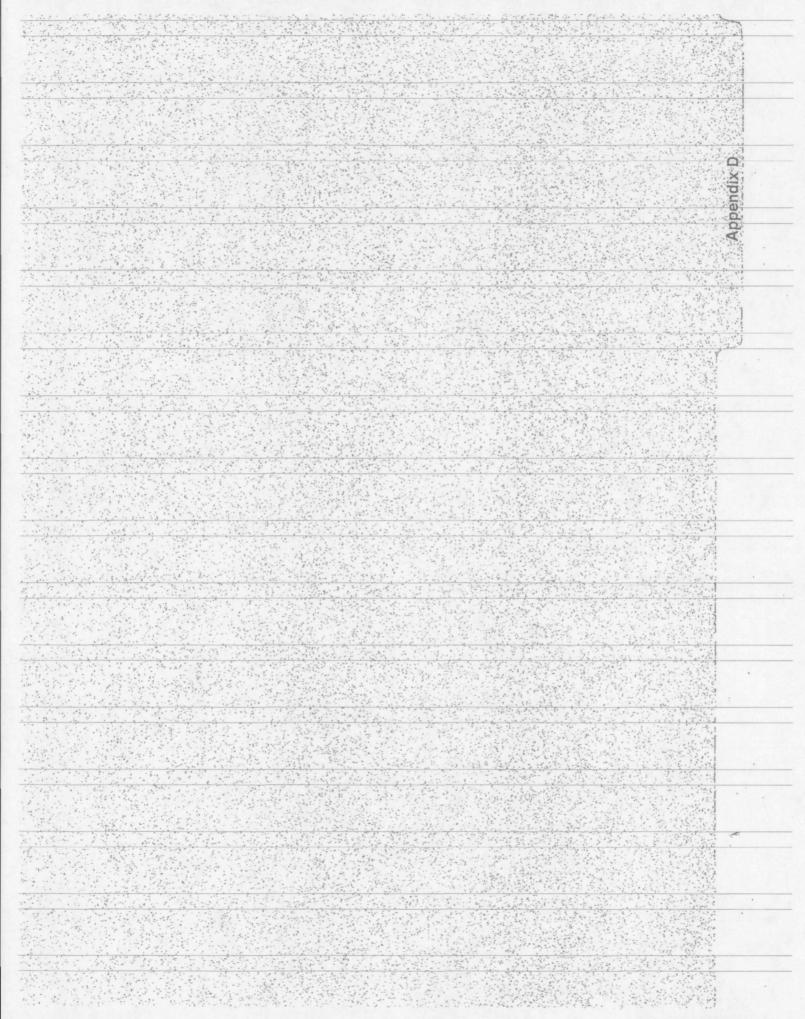
Photograph 12 - Facing north viewing south building on south side of Terminal 1.

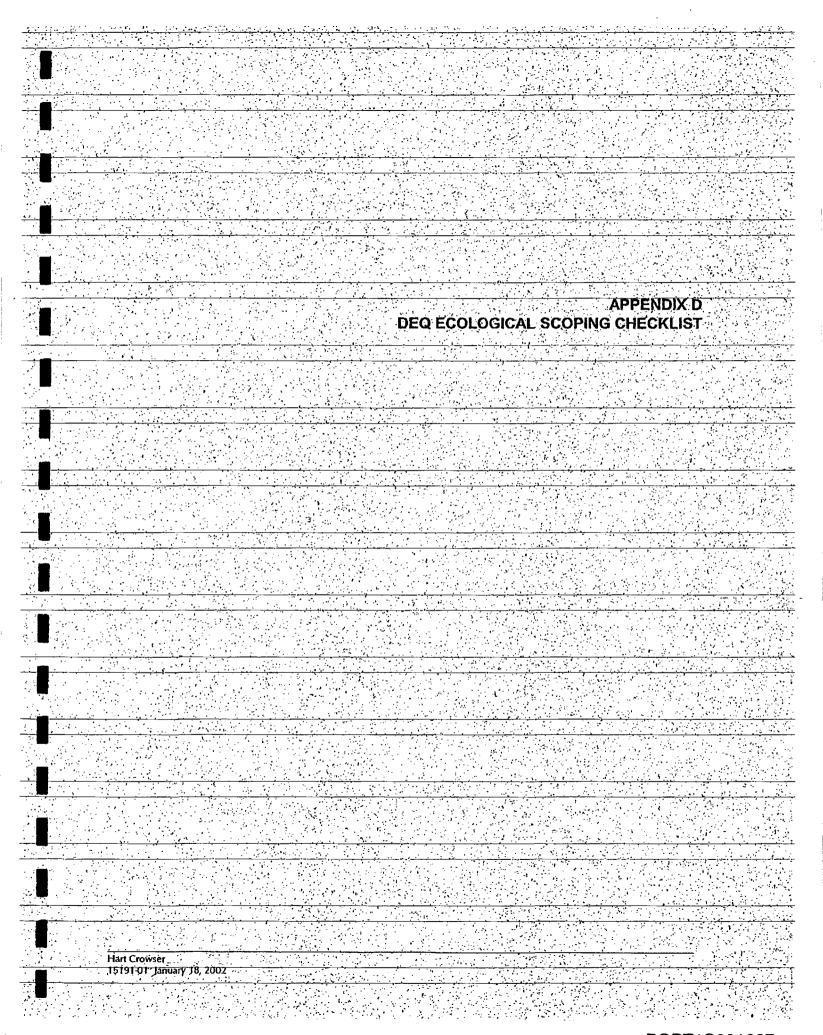


Photograph 13 -Facing south viewing south side of Terminal 1.



Photograph 14 -





ATTACHMENT 1 Ecological Scoping Checklist

Site Name	Terminal 1 South
Date of Site Visit	October 2, 2001
Site Location	2100 NW Front Avenue along the Willamette River in Portland, Oregon
Site Visit Conducted	Keith A. Kroeger
by	

Part 0

	CONTAMINANTS OF INTEREST Types, Classes, Or Specific Hazardous Substances †	Omete	Adjacent to or in locality of the facility †	
_	Known Or Suspected	Onsite	the facility	_
	Total Petroleum Hydrocarbons (TPH), polycyclic aromatic hydrocarbons	X		L
	(PAHs), volatile organic compounds (VOCs), and metals.			ŀ
1				
	•			Г
				
_				Н

¹ As defined by OAR 340-122-115(30)

Part @

_	OBSERVED IMPACTS ASSOCIATED WITH THE SITE	Finding	┞
	Onsite vegetation (None, Limited, Extensive)	N	I
	Vegetation in the locality of the site (None, Limited, Extensive)	N	
. •	Onsite wildlife such as macroinvertebrates, reptiles, amphibians, birds, mammals, other	N	Г
	(None, Limited, Extensive)		Γ
	Wildlife such as macroinvertebrates, reptiles, amphibians, birds, mammals, other in the	N	
	locality of the site (None, Limited, Extensive)		L
	Other readily observable impacts (None, Discuss below)	N	L
	Discussion:		
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[†] As defined by OAR 340-122-115(34)

ATTACHMENT 1 Ecological Scoping Checklist (cont'd)

SPECIFIC EVALUATION OF ECOLOGICAL RECEPTORS / HABITAT	Finding
Terrestrial - Wooded	
Percentage of site that is wooded	0
Dominant vegetation type (Evergreen, Deciduous, Mixed)	P *
Prominent tree size at breast height, i.e., four feet (<6", 6" to 12", >12")	
Evidence / observation of wildlife (Macroinvertebrates, Reptiles, Amphibians, Birds,	
Mammals, Other)	
Terrestrial - Scrub/Shrub/Grasses	. 1 3 2 3 1 1 1
Percentage of site that is scrub/shrub	<1%
Dominant vegetation type (Scrub, Shrub, Grasses, Other)	Weed
Prominent height of vegetation (<2', 2' to 5', >5')	<2'
Density of vegetation (Dense, Patchy, Sparse)	S
Evidence / observation of wildlife (Macroinvertebrates, Reptiles, Amphibians, Birds,	В
Mammals, Other)	
Terrestrial - Ruderal	
Percentage of site that is ruderal	99%
Dominant vegetation type (Landscaped, Agriculture, Bare ground)	B (paved)
Prominent height of vegetation (0', >0' to <2', 2' to 5', >5')	0'
Density of vegetation (Dense, Patchy, Sparse)	S
Evidence / observation of wildlife (Macroinvertebrates, Reptiles, Amphibians, Birds,	В
Mammals, Other)	
Aquatic - Non-flowing (lentic)	
Percentage of site that is covered by lakes or ponds	0%
Type of water bodies (Lakes, Ponds, Vernal pools, Impoundments, Lagoon, Reservoir,	
Canal)	
Size (acres), average depth (feet), trophic status of water bodies	
Source water (River, Stream, Groundwater, Industrial discharge, Surface water runoff)	
Water discharge point (None, River, Stream, Groundwater, Wetlands impoundment)	
Nature of bottom (Muddy, Rocky, Sand, Concrete, Other)	
Vegetation present (Submerged, Emergent, Floating)	
Obvious wetlands present (Yes / No)	
Evidence / observation of wildlife (Macroinvertebrates, Reptiles, Amphibians, Birds,	
Mammals, Other)	
Aquatic - Flowing (lotic)	
Percentage of site that is covered by rivers, streams (brooks, creeks), intermittent	0%
streams, dry wash, arroyo, ditches, or channel waterway	
Type of water bodies (Rivers, Streams, Intermittent Streams, Dry wash, Arroyo,	
Ditches, Channel waterway)	
Size (acres), average depth (feet), approximate flow rate (cfs) of water bodies	
Bank environment (cover: Vegetated, Bare / slope: Steep, Gradual / height (in feet))	
Source water (River, Stream, Groundwater, Industrial discharge, Surface water runoff)	
Tidal influence (Yes / No)	
Water discharge point (None, River, Stream, Groundwater, Wetlands impoundment)	
Nature of bottom (Muddy, Rocky, Sand, Concrete, Other)	

SPECIFIC EVALUATION OF ECOLOGICAL RECEPTORS / HABITAT	Finding
Vegetation present (Submerged, Emergent, Floating)	
Obvious wetlands present (Yes / No)	
Evidence / observation of wildlife (Macroinvertebrates, Reptiles, Amphibians, Birds,	
Mammals, Other)	
-Aquatic - Wetlands	
Obvious or designated wetlands present (Yes / No)	N
Wetlands suspected as site is/has (Adjacent to water body, in Floodplain, Standing	
water, Dark wet soils, Mud cracks, Debris line, Water marks)	
Vegetation present (Submerged, Emergent, Scrub/shrub, Wooded)	
Size (acres) and depth (feet) of suspected wetlands	
Source water (River, Stream, Groundwater, Industrial discharge, Surface water runoff)	
Water discharge point (None, River, Stream, Groundwater, Impoundment)	
Tidal influence (Yes / No)	
Evidence / observation of wildlife (Macroinvertebrates, Reptiles, Amphibians, Birds,	
Mammals, Other)	

^{*} P: Photographic documentation of these features is highly recommended.

Part @

	ECOLOGICALLY IMPORTANT SPECIES / HABITATS OBSERVED
	· · · · · · · · · · · · · · · · · · ·
	Terminal I South is located adjacent to the Willamette River, which is habitat for Coastal cutthroat trout (proposed threatened), Coho salmon (candidate), Steelhead (threatened), and Chinook salmon (threatened).
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ATTACHMENT 2 **Evaluation of Receptor-Pathway Interactions**

EVALUATION OF RECEPTOR-PATHWAY INTERACTIONS	Y	N	Ū	
Are hazardous substances present or potentially present in surface waters? AND		N		
Are ecologically important species or habitats present?	Y			
AND				
Could hazardous substances reach these receptors via surface water?		N		
When answering the above questions, consider the following:	<u> </u>			
Known or suspected presence of hazardous substances in surface waters.				
 Ability of hazardous substances to migrate to surface waters. 				•
• Terrestrial organisms may be dermally exposed to water-borne contaminants as a				
result of wading or swimming in contaminated waters. Aquatic receptors may be	1			
exposed through osmotic exchange, respiration or ventilation of surface waters.				
 Contaminants may be taken-up by terrestrial plants whose roots are in contact with surface waters. 		ì		
• Terrestrial receptors may ingest water-borne contaminants if contaminated surface				
waters are used as a drinking water source.				
Are hazardous substances present or potentially present in groundwater? AND		N		
Are ecologically important species or habitats present?	L _Y	<u> </u>		-
AND	_			
Could hazardous substances reach these receptors via groundwater?		N		
When answering the above questions, consider the following:				
Known or suspected presence of hazardous substances in groundwater.		\vdash		
Ability of hazardous substances to migrate to groundwater.				
• Potential for hazardous substances to migrate via groundwater and discharge into				
habitats and/or surface waters.				
• Contaminants may be taken-up by terrestrial and rooted aquatic plants whose roots are	1			
in contact with groundwater present within the root zone (~1m depth).				
• Terrestrial wildlife receptors generally will not contact groundwater unless it is				
discharged to the surface.				

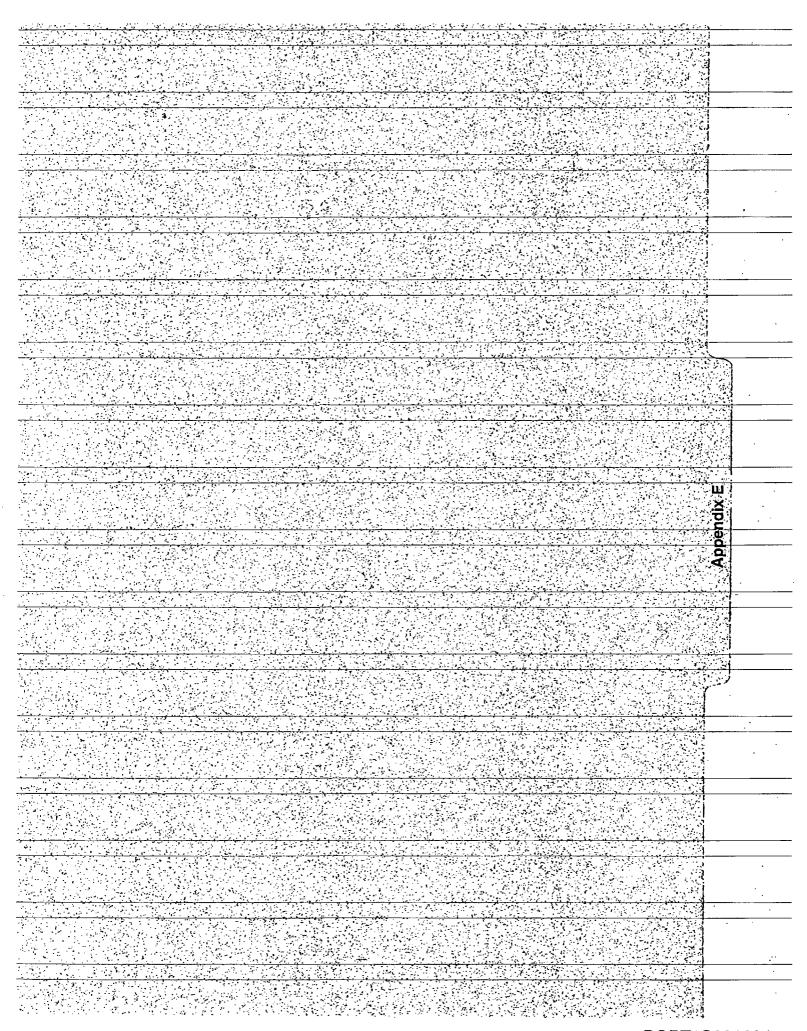
ATTACHMENT 2 Evaluation of Receptor-Pathway Interactions (cont'd)

EVALUATION OF RECEPTOR-PATHWAY INTERACTIONS	Y	N	U	
Are hazardous substances present or potentially present in sediments? AND		N		
Are ecologically important species or habitats present?	Y			
AND				
Could hazardous substances reach these receptors via contact with sediments?	ļ	N	ł	
When answering the above questions, consider the following:				
Known or suspected presence of hazardous substances in sediment.	<u> </u>			. :
 Ability of hazardous substances to leach or erode from surface soils and be carried 		Ι		
into sediment via surface runoff.				
Potential for contaminated groundwater to upwell through, and deposit contaminants	.			
in, sediments.	<u> </u>	_	\vdash	
If sediments are present in an area that is only periodically inundated with water,			,	
	ŀ			
terrestrial species may be dermally exposed during dry periods. Aquatic receptors	 	<u> </u>		
may be directly exposed to sediments or may be exposed through osmotic exchange,	+	-		
respiration or ventilation of sediment pore waters.	1 -			
• Terrestrial plants may be exposed to sediment in an area that is only periodically				
inundated with water.		 		
• If sediments are present in an area-that is only periodically inundated with water,	 	 		
terrestrial species may have direct access to sediments for the purposes of incidental		1	ŀ	
ingestion. Aquatic receptors may regularly or incidentally ingest sediment while		ļ	ll	
foraging.	!			
Are hazardous substances present or potentially present in prey or food items of	1	N		
ecologically important receptors?	1			
AND	_		l	
Are ecologically important species or habitats present?	Y			
AND				٠.
Could hazardous substances reach these receptors via consumption of food items?		N		
When answering the above questions, consider the following:				
 Higher trophic level terrestrial and aquatic consumers and predators may be exposed 	1		· · ·	
through consumption of contaminated food sources.				
• In general, organic contaminants with log $K_{ow} > 3.5$ may accumulate in terrestrial		1		
mammals and those with a log K _{ow} > 5 may accumulate in aquatic vertebrates.	1.	<u> </u>		

ATTACHMENT 2 Evaluation of Receptor-Pathway Interactions (cont'd)

EVALUATION OF RECEPTOR-PATHWAY INTERACTIONS	Y	N	U	
Are hazardous substances present or potentially present in surficial soils?	Υ			
AND				
Are ecologically important species or habitats present?		N		
ND		1		
Could hazardous substances reach these receptors via incidental ingestion of or		N		
lermal contact with surficial soils?	<u> </u>	L_		
When answering the above questions, consider the following:				
Known or suspected presence of hazardous substances in surficial (~1m depth) soils.				
Ability of hazardous substances to migrate to surficial soils.				
Significant exposure via dermal contact would generally be limited to organic	-	_		
contaminants which are lipophilic and can cross epidermal barriers.	-	-	·-· .	
Exposure of terrestrial plants to contaminants present in particulates deposited on leaf				
and stem surfaces by rain striking contaminated soils (i.e., rain splash).				
Contaminants in bulk soil may partition into soil solution, making them available to	1	\vdash	ļ-	
roots.			<u> </u>	
Incidental ingestion of contaminated soil could occur while animals grub for food	1			
resident in the soil, feed on plant matter covered with contaminated soil or while				
grooming themselves clean of soil.				
Are hazardous substances present or potentially present in soils?	Y			
AND	-			•
Are ecologically important species or habitats present?	<u> </u>	N	-	
AND	<u> </u>	<u> </u>		
Could hazardous substances reach these receptors via vapors or fugitive dust carried		N		
n surface air or confined in burrows?	Ì	l		
When answering the above questions, consider the following:	F			
Volatility of the hazardous substance (volatile chemicals generally have Henry's Law				
constant > 10 ⁻⁵ atm-m³/mol and molecular weight < 200 g/mol).				
Exposure via inhalation is most important to organisms that burrow in contaminated				٠.
soils, given the limited amounts of air present to dilute vapors and an absence of air				
movement to disperse gases.				
• Exposure via inhalation of fugitive dust is particularly applicable to ground-dwelling		ľ		
species that could be exposed to dust disturbed by their foraging or burrowing	<u> </u>	L		
activities or by wind movement,		<u> </u>	<u> </u>	
Foliar uptake of organic vapors would be limited to those contaminants with relatively				
high vapor pressures.		1		
Exposure of terrestrial plants to contaminants present in particulates deposited on leaf-	<u> </u>	<u> </u>		
and stem surfaces.		<u> </u>	\vdash	
1/22 (SN22				1

"Y" = yes; "N" = No, "U" = Unknown (counts as a "Y")

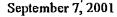


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NATURAL HERITAGE PROGRAM

A Cooperative Project of:







Taku Fuji, Ph.D.
Hart Crowser, Inc.
Five Centerpointe Drive, Suite 240
Lake Oswego, OR 97035-8652

1322 SE Morrison Street Portland, OR 97214-2423 VOICE/FAX (503) 731-3070

Dear Dr. Fuji:

Thank you for requesting information from the Oregon Natural Heritage Program (ONHP). We have conducted a data system search for rare, threatened and endangered plant and animal records for your Port of Portland Terminal 1 South Project (15191-01) in Township 1 North, Range 1 East, Section 28.

Fourteen (14) records were noted within a two-mile radius of your project and are included on the enclosed computer printout. A key to the fields is also included.

Please remember that the lack of rare element information from a given area does not mean that there are no significant elements there, only that there is no information known to us from the site. To assure that there are no important elements present, you should inventory the site, at the appropriate season.

Please note that at this time ONHP does not have comprehensive computerized records available for all anadromous fish in Oregon. I have listed below the species that may be present within the waterways contained in the project area. I have also included their listing by the National Marine Fisheries Service (NMFS). For more information on anadromous fish you may wish to contact NMFS at: 525 NE Oregon Street; Portland, Oregon 97232-2737. Please also note that the U.S. Fish and Wildlife Service now has jurisdiction over coastal cutthroat trout.

Coastal-cuttlroat-trout-Oncorhynchus clarki-clarki-Proposed Threatened (Columbia River/SW Washington) Coho salmon (Lower Columbia River) Oncorhynchus kisutch Candidate Steelhead (Lower Columbia River) Oncorhynchus mykiss Threatened Oncorhynchus mykiss Threatened Steelhead (Upper Willamette River) Chinook salmon (Lower Columbia River) Oncorhynchus tshawytscha Threatened Chinook salmon (Upper Willamette River) Oncorhynchus tshawytscha Threatened

This data is confidential and for the specific purposes of your project and is not to be distributed.

If you need additional information or have any questions, please do not hesitate to contact me.

Sincerely,

Cliff Alton

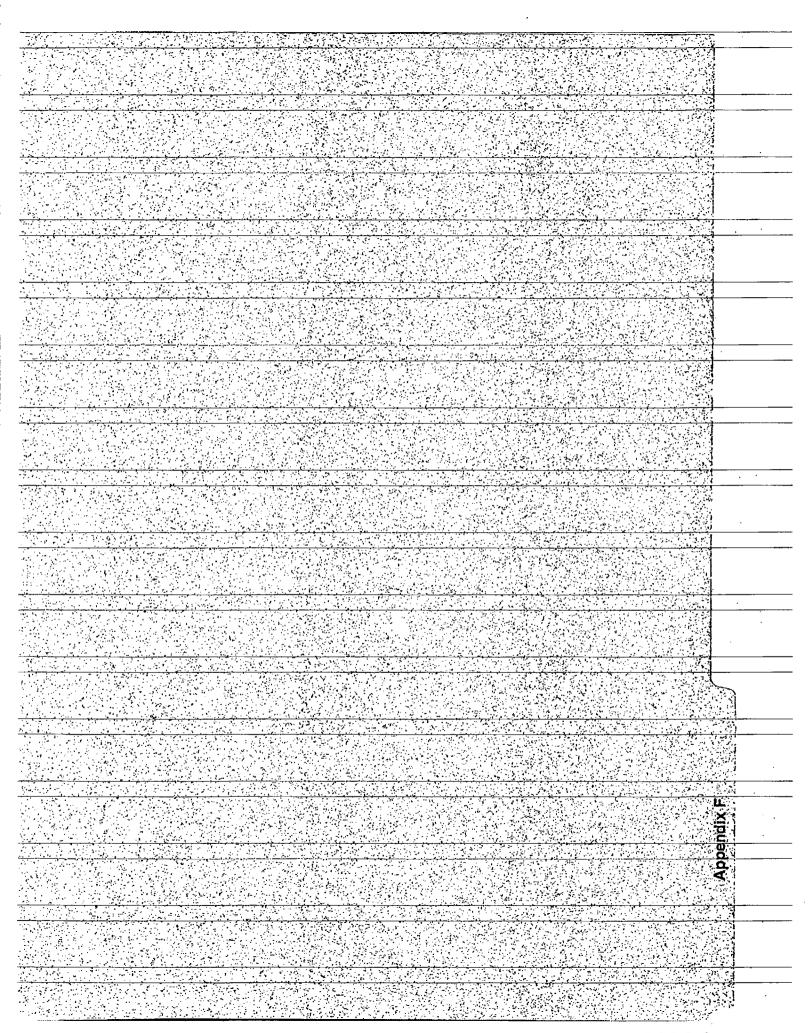
Conservation Information Assistant

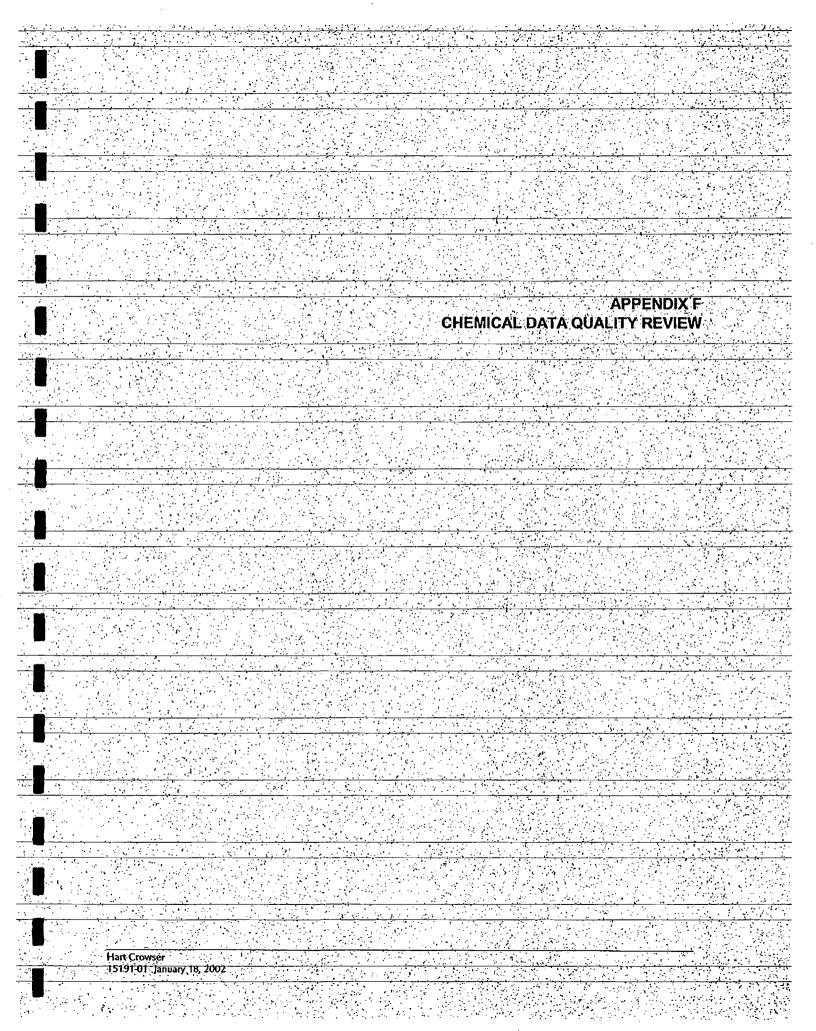
computer printout and data key

From: Uliff Aifon - 505-751-3	07 0-10:-13KU FUJI	Nate: Attivior i	ma; 0;20;40-PM	rage 3-01-1U
•				
18:18:04 07 SEP 2001		Page 2		
			·	
MAME.	ONCORHYNCHUS TSHAMYTSCHA		•	
	CHINOOK SALMON - LOWER C			
E0-GODE:-	-AFCHA0205W*006	LAST_088:-1999-PRE	FED-STATUS: LT	
COUNTY(8):		FIRST OBS:	STATE STATUS:	•
	MULTNOMAH . COLUMBIA	•		
QUAD NAMES:	OREGON CITY	LAT:	SIZE:	
	GLADSTONE			
=	LAKE OSWEGO			
	PORTLAND			
,	-LINNTON			
	ST HELENS			
PHYSIOGRAPHIC PROV:		LONG:	MINELEV (Feet):	
T-R-S:		QUADCODE: 4512235	MAXELEV (Feet):	
·	***	45 12245 45 12246		
		4512256	•	•
	•	4512257	•	•
		4512267		
		4512277		
T-R-S COMMENTS: EO-RANK/COMM:			PRECISION: H	
		CHANNEL, WILLAMETTE RIVER		
DESCRIPTION:				
		UTION NAPS USED TO CREATE THE 1:	24,000 COVERAGE	
	REARING & MIGRATION - fi	·	OM ODFW GEOGRAPHIC RESOURCES DATA	PRODUCER AND
			DATA FIELD, THE INFORMATION PRESEN	
J .			STRICT FISHERIES BIOLOGIST; THE PR	
		AS SHOULD BE CONSIDERED UNDOCUME	NTED BUT AS HAVING A POTENTIAL OF	DEING PRESENT.
ANNUAL OBSERVATION:				•
MANAGED AREA:			<u> </u>	
MANAGE COMM:	·	·		
PROT COMM:	•	·	•	
BEST SOURCE:	2000 ODFN GEOGRAPHIC RES	OURCES DATA; MASSEY, JAY; BENNET	T, DON.	
NAME:	ONCORHYNCHUS TSHANYTSCHA	POP 22		
	CHINOOK SALHON - LOWER C	•		
—	AFCH40205Y*006	LAST 08S: 1999-PRE	FED STATUS: LT	
COUNTY(B):		FIRST OBS:	STATE STATUS: SC	
	MULTNOMAH COLUMBIA			
QUAD NAMES:	OREGON CITY	LAT:	SIZE:	.
	GLADSTONE			
_======================================	LAKE OSHEGO			•
	PORTLAND			
-	SAUVIE ISLAND			
PHYSIOGRAPHIC PROV:	5.5,15	LONG;	MINELEV (Feet):	
		QUADCODE: 4512235	MAXELEV (Feet):	
		4512245		
	•	* 4512240 4512256		
		4512256 4512257		•
		45122 67		
T-R-S CONMENTS:		· · · · · · · · · · · · · · · · · · ·	PRECISION: H	
EO-RANK/COM:				
DIRECTIONS: DESCRIPTION:	SCAPPOOSE BAY & TRIBUTAR	IES, WILLAMETTE RIVER & TRIBUTAR	162	
DEGUNTE 140W.				

FIUNE UNITABLE DUSTES ISSUED IN LAND FUR 8:19:04 07 SEP 2001 Page 3 EO-DATA: FALL RUN; ODFN DISTRIBUTION MAPS USED TO CREATE THE 1:24,000 COVERAGE EOTYPE: REARING & MIGRATION - fish COMMENTS: DISTRIBUTION INFORMATION USED IN THIS EOR WAS DERIVED FROM ODEN GEOGRAPHIC RESOURCES DATA PRODUCED AND DISTRIBUTED_IN_1999._UNLESS_SPECIFIC_DATA_EXISTS_IN_THE_DATA_FIELD,_THE_INFORMATION_PRESENTED_IN_THIS_EOR. REPRESENTS THE "BEST PROFESSIONAL JUDGMENT" BY ODFH'S DISTRICT FISHERIES BIOLOGIST; THE PRESENCE OF CHINOOK IN DESCRIBED AREAS SHOULD BE CONSIDERED UNDOCUMENTED BUT AS HAVING A POTENTIAL OF BEING PRESENT. ANNUAL OBSERVATION: OHNER: HANAGED AREA: HANAGE COHM: PROT COMM: BEST SOURCE: 2000 ODEN GEOGRAPHIC RESOURCES DATA; MASSEY, JAY; BENNETT, DON; CALDNELL, DICK. NAME: ONCORHYNCHUS MYKISS POP 27 COMMON NAME: STEELHEAD . LOWER COLUMBIA RIVER WINTER RUN EO-CODE: AFGHA02132*001 LAST 088: 1999-PRE FED STATUS: LT COUNTY(8): CLACKAMAS FIRST ORS: BTATE STATUS: SC HANONT-JUH COLUXBIA QUAD NAMES: OREGON CITY LAT: SIZE: GLADSTONE LAKE OSWEGO PORTLAND LINATON SAUVIE ISLAND ST HELENS HYSIOGRAPHIC_PROV LONG: MINELEV (Foot): QUADCODE: 4512235 MAXELEV (Feet): 4512245 4512246 4512256 4512257 4512267 4512277 T-R-8 COMMENTS: PRECISION: M EO-RANK/COMM: DIRECTIONS: SCAPPOOSE BAY, MULTNOHAH CHANNEL, MILLAMETTE RIVER EO-DATA: WINTER RUN: ODFN DISTRIBUTIION MAPS USED TO CREATE THE 1:24,000 COVERAGE EUTYPE: REARING & MIGRATION - fish COMMENTS: DISTRIBUTION INFORMATION USED IN THIS EOR WAS DERIVED FROM ODEW GEOGRAPHIC RESOURCES DATA PRODUCED AND DISTRIBUTED IN 1999. UNLESS SPECIFIC DATA EXISTS IN THE DATA FIELD, THE INFORMATION PRESENTED IN THIS EOR REPRESENTS THE *BEST PROFESSIONAL JUDGMENT* BY ODFH'S DISTRICT FISHERIES BIOLOGIST; THE PRESENCE OF STEELHEAD IN DESCRIBED AREAS SHOULD BE CONSIDERED UNDOCUMENTED BUT AS HAVING A POTENTIAL OF BEING AMMUAL OBSERVATION: OWNER: MANAGED AREA: MANAGE COMM: BEST SOURCE: 2000 ODFW GEOGRAPHIC RESOURCES DATA; MASSEY, JAY; BENNETT, DON. NAME: CORYNORHINUS TOWNSENDII TOWNSENDII COMMON NAME: PACIFIC WESTERN BIG-EARED BAT EO-CODE: AMACCO8015*071 LAST 085: 1928-09-05 FED STATUS: SOC COUNTY(s): MULTHOMAH FIRST 08S: 1914 STATE STATUS: SC QUAD NAMES: PORTLAND LAT: 453220N SIZE: 0 PHYSIOGRAPHIC-PROV: MV LONG: 1223800H MINELEV (Feet): 150 T-R-S: 00111001E 25 QUADCODE: 4512256 HAXELEV_(Feet): T-R-S.COMMENTS: PRECISION: G EO-RANK/COMM: D DIRECTIONS: PORTLAND - ON THE E SIDE

FIRST CHILDREN SWITT STITUTE 10. TORGET OF 18:18:05 07 SEP 2001 Page 5 T-R-S COMMENTS: NWA [TRS NOT GIVEN] PRECISION: M EO-RANK/COMM: D : DIRECTIONS:-HOYT PARK, FAIRVIEW BOULEVARD. DESCRIPTION: EO-DATA: 1965: 1 INDIVIDUAL COLLECTED **EOTYPE:** CONHENTS: OBSERVER: CAVANAGH, R. PORTLAND STATE UNIVERSITY SPECIMEN #002431. ANNUAL OBSERVATION: · OWNER: CITY MANAGED AREA: HANAGE COMM: PROT COMM: BEST SOURCE: BRUCE, CHARLIE. ODFW. NAME: CLEMMYS MARNORATA MARMORATA COMMON NAME: NORTHWESTERN POND TURTLE E0-C0DE: ARAAD02031*041 FED-STATUS: SOC EAST OBS: FIRST OBS: STATE-STATUS:-SC COUNTY(&): MULTHOMAH -QUAD NAMES: PORTLAND LAT: 453045N SIZE: 0 PHYSIOGRAPHIC PROV: NV LONG: 1224130M MINELEV (Feet): T-R-S: 0015001E QUADCODE: 4512258 MAXELEV (Feet): PRECISION: G T-R-S COMMENTS: EO-RANK/COMM: D DIRECTIONS: PORTLAND -EO-DATA: SPECIES RECORDED AT THIS-SITE, PER ST. JOHN. DATES NOT SPECIFIED EOTYPE: COMMENTS: ANNUAL OBSERVATION: OHNER: MANAGED AREA: HANAGE COMH: PROT COMM: BEST SOURCE: ST. JOHN, ALAN: 1984. HERPETOLOGY OF THE LOWER WILLAMETTE VALLEY NAME: ASTER CURTUS COMMON NAME: WHITE-TOPPED ASTER LAST 08S: 1898-08 FED STATUS: SOC EO-CODE: PDASTOTORO*008 FIRST 085: 1898 STATE STATUS: LT COUNTY(s): HULTNONAH QUAD NAMES: PORTLAND LAT: 453045N SIZE: 0 LONG: 1224130H -MINELEV (Feet): 100-PHYSIOGRAPHIC PROV: HV T-R-S: 0015001E QUADCODE: 4512256 MAXELEV (Feet): PRECISION: G T-R-S COMMENTS: EO-RANK/COMM: DIRECTIONS: PORTLAND. NO FURTHER INFORMATION. DESCRIPTION: NO HABITAT DATA. EO-DATA: HERBARIUN COLLECTION: NO NAME, 08-09-98, NO #, WS -COMMENTS: THIS POPULATION ASSUMED EXTINPATED. ANNUAL_OBSERVATION: CHNER: MANAGED AREA: MANAGE COMM: PROT COMM: BEST SOURCE: HERBARIUM COLLECTION AT HS NAME: CINICIFUGA ELATA COMMON NAME: TALL BUGBANE LAST 08S: 1904-05-30 FED STATUS: EO-CODE: PORANO7030*017 FIRST 089: 1882 STATE STATUS: C COUNTY(8): MULTNOHAH QUAD NAMES: PORTLAND LAT: 453045N SIZE: PHYSIOGRAPHIC PROV: WV LONG: 1224130H **HINELEV** (Feet):





APPENDIX	
	DATA QUALITY REVIEW ORTLAND- TERMINAL 1
PORTLAND	
	Hahn and Associates, Inc. (HAI) of Portland, Oregon, submitted soil and
	water samples to various laboratories for analysis in 1998, 2000, and 2001.
	The laboratories included Oregon Analytical Laboratory (OAL) of Beaverton,
	Oregon; North Creek Analytical, Inc. (NCA) of Portland, Oregon; and
	Environmental Services Laboratory, Inc. (ESL) of Portland, Oregon. Hart
	Crowser has performed cursory reviews of laboratory data compiled by HAI
	in Volumes 1 and 2 of a document titled "Terminal 1 South Remedial
	Investigation Report" and the "Monitoring Well Installation and Groundwater Sampling Report".
	The following criteria were evaluated in the data quality review process:
	■ Holding times;
	■ Method blanks;
	■ Surrogate recoveries;
	 Laboratory control sample/laboratory control sample duplicate (LCS/LCSD) recoveries;
	■ Matrix spike/matrix spike duplicate (MS/MSD) recoveries; and
	 Laboratory and field duplicate relative percent difference (RPD).
·	The review is organized by the Appendix containing the data, and is further
	subdivided by laboratory.
Appendix A	: Focused Environmental Site Assessment (August 1998)
	NCA ID No. P803593
	Twenty-two soil samples were submitted to the laboratory. The following
	analyses were performed on one or more samples:
	 Hydrocarbon Identification (modified EPA Method 8015);
	■ Total Metals (EPA 6000/7000 Series Methods);
	Phenols (EPA Method 8040A);

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	TO STATE OF THE ST	
	Organochlorine Pesticides (EPA Method 8081);	
	■ Polychlorinated Biphenyls (PCBs, EPA Method 8081);	•
	■ Volatile Organic Compounds (VOCs, EPA Method 8260A);	
	■ Chlorinated Herbicides (EPA Method 8151A); and	
•		
	■ Total Petroleum Hydrocarbon (TPH) as Diesel/Oil (NWTPH-Dx).	
	Hydrocarbon Identification. All required holding times were met. No	
	method blank contamination was detected. Surrogate recoveries were	
	acceptable. No duplicate results were provided.	
	Total Metals. All required holding times were met. No method blank	
	contamination was detected. LCS and MS recoveries were within laboratory	
	control limits with the following exception from Batch 0380703. Antimony	
· · · · · · · · · · · · · · · · · · ·	MS recovery 26 percent was less than control limits (75 to 125 percent),	
	associated antimony results, which were all nondetect, were flagged as	
	estimated ("U)"). Laboratory duplicate RPDs were acceptable with the	
	following exceptions for Batch 0380703. Selenium was detected in the	,
	laboratory duplicate, but was not detected in the original sample. However,	
	since the laboratory duplicate result is less than five times the sample	
	reporting limit, no qualification was necessary. The duplicate RPD for arsenic	
	(68.2 percent) exceeded the control limit of 40 percent. Associated arsenic	
	results were flagged as estimated ("J"). No field duplicates were identified.	
	Phenols. All required holding times were met. No method blank	
	contamination was detected. Surrogate and LCS recoveries were within	
	laboratory control limits with the following exceptions. The LCS recoveries	
 	for 4-chloro-3-methylphenol, 2-chlorophenol, and phenol were greater than	
	control limits, while the LCS duplicate recovery for 4-nitrophenol was greater	
	than the control limit. The LCS duplicate RPDs were acceptable. Since no	
	phenolic compounds were detected, no qualifiers were necessary. No field	
·	duplicates were identified.	
	Organochlorine Pesticides. All required holding times were met. No	
	method blank contamination was detected. Surrogate, LCS, and MS	
	recoveries were within laboratory control limits. MS/MSD RPDs were not	
<u> </u>	calculated since the MSD sample extract was lost during the GPC procedure.	
	No field duplicates were identified.	
•	PCBs. All required holding times were met. No method blank contamination	
	was detected. Surrogate, LCS, and MS recoveries were within laboratory control	
	limits. MS/MSD RPDs were acceptable. No field duplicates were identified.	
· · · · · · · · · · · · · · · · · · ·		•
Hart Crowser	Page F-2	

· .	
•	VOCs. All required holding times were met. No method blank
	contamination was detected. Surrogate, LCS, and MS recoveries were within
	laboratory control limits. MS/MSD RPDs were acceptable. No field
	duplicates were identified.
	Chlorinated Herbicides. All required holding times were met. No method
	blank contamination was detected. Surrogate and LCS recoveries were
	within laboratory control limits. LCS duplicate RPDs were acceptable. No
	field duplicates were identified.
	TPH as Diesel/Oil. All required holding times were met. No method blank
	contamination was detected. Surrogate and LCS recoveries were within
· .	laboratory control limits with the following exception. The surrogate recovery
	for sample T-1 B-5 0-2 was not recoverable due to sample dilution. No
	qualification was necessary. Laboratory duplicate RPDs were acceptable. No
	field duplicates were identified.
A	In Depoling Spill Sounday (Fabruage and March 2000)
Appenaix	I: Baseline Soil Samples (February and March 2000)
	Sample No. Prefix: 4976-000229
•	OAL ID No. L153236
	Twenty-nine soil samples were submitted to the laboratory. The following
	Twenty-nine son samples were submitted to the laboratory. The following
	analyses were performed on one or more samples:
	analyses were performed on one or more samples:
	·
	analyses were performed on one or more samples:
	analyses were performed on one or more samples: Total Metals (EPA 6000/7000 Series Methods);
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260);
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID);
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260);
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID);
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082);
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx).
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory control limits with the following exception. Antimony MS recovery was 72
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory control limits with the following exception. Antimony MS recovery was 72 percent versus the control limit of 75 to 125 percent. Associated antimony
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory control limits with the following exception. Antimony MS recovery was 72 percent versus the control limit of 75 to 125 percent. Associated antimony results, which were-all-nondetect, were flagged "UJ". The MS recovery for
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory control limits with the following exception. Antimony MS recovery was 72 percent versus the control limit of 75 to 125 percent. Associated antimony results, which were all nondetect, were flagged "UJ". The MS recovery for lead was less than the control limit for sample -020; however, since the spike
	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory control limits with the following exception. Antimony MS recovery was 72 percent versus the control limit of 75 to 125 percent. Associated antimony results, which were-all-nondetect, were flagged "UJ". The MS recovery for
Hart Crowser	analyses were performed on one or more samples: ■ Total Metals (EPA 6000/7000 Series Methods); ■ Benzene, Toluene, Ethylbenzene, and Xylene (BTEX, EPA Method 8260); ■ Volatile Organic Compounds (VOCs, EPA Method 8260); ■ TPH Identification (NWTPH-HCID); ■ PCBs (EPA Method 8082); ■ Polynuclear Aromatic Hydrocarbons (PAHs, EPA Method 8270 SIM); and ■ TPH as Diesel/Oil (NWTPH-Dx). Total Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory control limits with the following exception. Antimony MS recovery was 72 percent versus the control limit of 75 to 125 percent. Associated antimony results, which were all nondetect, were flagged "UJ". The MS recovery for lead was less than the control limit for sample -020; however, since the spike

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	· · · · · · · · · · · · · · · · · · ·
	magazane Labarratan dunligata PPDs ware acceptable. No field dunligator
	necessary. Laboratory duplicate RPDs were acceptable. No field duplicates
	were identified.
	BTEX. All required holding times were met. No method blank contamination
	was detected. Surrogate, LCS, and MS recoveries were within laboratory
	control limits. MS/MSD duplicate RPDs were acceptable. The MS/MSD data
	reported for sample -020 were prepared/analyzed six days before the sample
	was prepped, and is therefore unacceptable. Data for sample -020 was
	rejected. No field duplicates-were identified.
	VOCs. All required holding times were met. No method blank
	contamination was detected. Surrogate, LCS, and MS recoveries were within
	laboratory control limits with the following exception. The toluene-d8
	recovery for sample -028 was greater than laboratory control limits.
	However, since this is a positive bias and all results were not detected, no
	qualifiers were necessary. No field duplicates were identified.
•	TD11 8 Jane 25 and an All required by a Jaimer Singer Course week. No most had blood
	TPH Identification. All required holding times were met. No method blank
	contamination-was detected. Surrogate recoveries were within laboratory
	control limits with the following exception. The 2-fluorobiphenol recovery for sample was not reported due to dilution. No qualifiers were necessary.
	Laboratory duplicate RPDs were acceptable. No field duplicates were identified.
	Laboratory auplicate in 155 were acceptable. 140 neta duplicates were identified.
	PCBs. All required holding times were met. No method blank contamination
	was detected. Surrogate, LCS, and MS recoveries were within laboratory
	control limits. MS/MSD RPDs were acceptable. The LCS and MS/MSD data
	reported for sample -020 was prepared/analyzed seven before the sample
	was prepped, and is therefore unacceptable. Data for sample -020 was
	rejected. No field duplicates were identified.
	PAHs. All required holding times were met with the exception of sample -
	020, which was analyzed 23 days after the sample was collected. All
	detected results were flagged "J", while nondetected results were flagged
	"UJ". No method blank contamination was detected. Surrogate, LCS, and
	MS recoveries were within laboratory control limits. MS/MSD RPDs were
	acceptable. No field duplicates were identified.
	TPH-as-Diesel/Oil. All required holding times were met. No method blank
	contamination was detected. Surrogate, LCS, and MS recoveries were within
	laboratory control limits with the following exception. The surrogate recovery
	for sample -020 was not recoverable due to sample dilution. No qualification
	was necessary. Laboratory duplicate RPDs were acceptable. The LCS and
	MS data reported for samples -003, -011, -012, -013, -014, -020, -021, -023,
	1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1
Hart Crowser	Page F-4

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	and -028 were prepared/analyzed between 1 and 2 weeks before the
	samples were prepped, and are therefore unacceptable. Data for these
	samples were rejected. No field duplicates were identified.
Appendi	ix J: B-38 Area Characterization (March-2000)
	Sample No. Prefix: 4876-000313 OAL ID No. L15469
	Thirty-one soil samples were submitted to the laboratory, including some
	samples that were not analyzed in this set. The following analyses were
	performed on one or more samples:
	■ TPH as Diesel/Oil (NWTPH-Dx);
	 Aromatic Hydrocarbons (BTEX, EPA Method 8020A); and
	■ PAHs (EPA Method 8270 SIM).
	TPH as Diesel/Oil. All required holding times were met. No method blank
	contamination was detected. Surrogate recoveries were generally within
	laboratory control limits. Surrogate recoveries in some soil samples were
	above control limits due to high sample concentrations. Duplicate, LCS, and
	MS data reported for some samples were analyzed several days before the
	samples arrived in the laboratory. TPH data for samples -031, -036, -051, and
	-056 were rejected on this basis. The remaining duplicate and LCS data were
	within control limits. MS recoveries for samples -043, -044, -49, -053, -058,
	and -060 were outside of control limits, but the LCS data were acceptable, so
·	no data were qualified on this-basisNo field duplicates were identified
	BTEX. All required holding times were met. No method blank contamination
	was detected. LCS, surrogate, and MS recoveries were within laboratory
	control limits. Laboratory duplicate RPDs were acceptable. No field
	duplicates were identified.
	PAHs. All required holding times were met. No method blank contamination
	was detected. LCS, surrogate, and MS/MSD recoveries were within
	laboratory control limits. Laboratory duplicate RPDs were acceptable. No
	field duplicates were identified.

,	
	·
	Sample Prefix No. 4876-000316
· ·	OAL ID No. L15520
	OAL 10 110. L13320
	Thirty soil samples were submitted to the laboratory, including some samples
	that were not analyzed in this set. The following analyses were performed on
	one or more samples:
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	■ TPH Identification (NWTPH-HCID);
	■ TPH as Diesel/Oil (NWTPH-Dx);
	■ BTEX, (EPA Method 8020A); and
	■ PAHs, (EPA Method 8270 SIM).
	· · · · · · · · · · · · · · · · · · ·
	TPH Identification. All required holding times were met. No method blank
	contamination was detected. LCS and MS analyses are not required by
•	NWTPH-HCID procedure. Surrogate recoveries were acceptable. The
	laboratory duplicate RPDs reported were extracted three days before the
	incumbent samples were extracted, and are therefore unacceptable. Samples
	-063 and -073 were flagged "J" based on the duplicate data and lack of
	suitable additional qualify control. No field duplicates were identified.
	TPH as Dicsel/Oil. All required holding times were met. No method blank
	contamination was detected. LCS and surrogate recoveries were within
	_
	laboratory control limits. MS spiking levels were less than 20 percent of the native soil concentrations; therefore the control limits do not apply.
	•••
	Laboratory duplicate RPDs were acceptable. Duplicate, LCS, and MS data
	reported for -082 and -084 were prepared the day before the sample was
	prepared, and is therefore unacceptable. Data for -082 and -084 were
	rejected. No field duplicates were identified.
	BTEX. All required holding times were met. No method blank
	contamination was detected. LCS, surrogate, and MS recoveries were within
	laboratory control limits. Laboratory duplicate RPDs were acceptable. No
	field duplicates were analyzed. Duplicate, LCS, and MS data reported for -
····	075 and 079 were prepared/analyzed three days before the sample was
	prepared, and is therefore unacceptable. Data for -075 and -079 were
	rejected. No field duplicates were identified.
***************************************	PAHs. All required holding times were met. No method blank
	contamination was detected. LCS/LCSD and surrogate recoveries were
 	within laboratory control limits. Laboratory duplicate RPDs were acceptable.
·	No field duplicates were analyzed. LCS/LCSD data reported for -075 and -

acceptable. No field duplicates were identified.
qualified as a result of the MS values. Laboratory duplicate RPDs were
limits; however, since-LCS-recoveries-were-acceptable, no-data-were-
laboratory control limits. MS recoveries for all analytes were below control
contamination was detected. LCS and surrogate recoveries were within
VOCs. All required holding times were met. No method blank
field duplicates were identified.
laboratory control limits. Laboratory duplicate RPDs were acceptable. No
was detected. LCS, surrogate, and MS/MSD recoveries were within
PAHs. All required holding times were met. No method blank contamination
were identified.
data were flagged as a result of the duplicate analyses. No field duplicates
laboratory duplicate RPDs was outside the laboratory control limits. No
laboratory control limits. No MS analyses were performed. One of two
contamination was detected. LCS and surrogate recoveries were within
TPH as Diesel/Oil. All required holding times were met. No method blank
■ Tributyltin (TBT, Krone method).
■ Priority Pollutant Metals (EPA 6000/7000 Series Methods); and
■ PCBs (EPA Method 8082);
■ Low-level VOCs (EPA Method 8260B);
■ PAHs (EPA Method 8270 SIM);
■ TPH as Diesel/Oil (NWTPH-Dx);
performed on one of more samples:
performed on one or more samples:
Forty-two soil samples were submitted to the laboratory, including some samples that were not analyzed in this set. The following analyses were
Sample Prefix Nos. 5106-000919 & 5106-000920 NCA ID No. P009611
x K: Supplemental Site Characterization (September 2000)
field duplicates were identified.
079 were prepared and analyzed five days before the sample was analyzed and is therefore unacceptable. Data for -075 and -079 were rejected. No
·

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	laboratory control limits. LCS analysis is not required by the method, and
	contamination was detected. Surrogate and MS recoveries were within
•	TPH as Diesel/Oil. All required holding times were met. No method blank
	TCLP Lead (EPA 1311/6000/7000 Methods).
•	
	■ Priority Pollutant Metals (EPA 6000/7000 Series Methods); and ^C
	■ Polychlorinated Biphenyls (PCBs, EPA Method 8082);
	■ VOCs (EPA Method 8260B);
	■ PAHs (EPA Method 8270 SIM);
· · · · · · · · · · · · · · · · · · ·	■ TPH as Diesel/Oil (NWTPH-Dx);
	■ TPH as Diocol/Oil /NIM/TPH DV).
	performed on one-or-more samples:
	samples_that-were-not-analyzed in this set. The following-analyses-were
	Thirty-six soil samples were submitted to the laboratory, including some
	NCA ID No. P009722
	Sample Prefix Nos. 5106-000919 & 5106-000920
· · · · · · · · · · · · · · · · · · ·	mormoo paii, and were acceptable. 140 field duplicates were identified.
	limits were reported. Duplicate analysis RPD data were limited to the MS/MSD pair, and were acceptable. No field duplicates were identified.
	performed. MS/MSD analyses were performed, but no established control
<u></u>	detected. Surrogate recoveries were acceptable. No LCS analysis was
	holding times could not be identified. No method blank contamination was
	reported using the PSEP protocols to perform the analysis. Established
	TBT. Analysis was subcontracted to Sound Analytical Services, which
	field duplicates were identified.
	recovery was acceptable. Laboratory duplicate RPDs were acceptable. No
	lead was slightly below control limits. Results were not qualified since the MS
	laboratory control limits with the following exceptions. MSD recovery for
	blank contamination was detected. LCS and MS recoveries were within
	Priority Pollutant Metals. All required holding times were met. No method
	and was acceptable. No field duplicates were identified.
	Laboratory duplicate RPD data were limited to LCS/LCSD and MS/MSD pairs
	MS, and surrogate recoveries were within laboratory control limits.
	expired holding time. No method blank contamination was detected. LCS,
	the persistent nature of the analyte, no data were qualified as a result of the
	023 was re-extracted 8 days after the holding time expired; however, given
	PCBs. All required holding times were met with one exception. Sample -
•	
· · · · · · · · · · · · · · · · · · ·	

none was performed. Laboratory duplicate RPDs were acceptable. No field duplicates were identified. PAHs. All required holding times were met. No method blank contamination was detected. LCS and surrogate recoveries were within laboratory control limits. MS/MSD results were within control limits with one exception. 0100976MSD1 had a pyrene recovery above the laboratory control limits; however, since the bias was positive and the LCS results were accepted, no data were qualified as a result of the high pyrene recovery. Laboratory duplicate RPDs for -050 were outside control limits. Given the acceptability of the balance of the PAH QC data, no qualifiers were assigned. No field duplicates were identified. **VOCs.** All required holding times were met. No method blank contamination was detected. LCS and surrogate recoveries were within laboratory control limits with one exception. The 4-BFB surrogate for -069 had a recovery above the upper control limit; however, since the bias was positive, no qualifiers were assigned. One MS/MSD set had recoveries for all analytes below control limits; however, since LCS recoveries were acceptable, no data were qualified as a result of the MS/MSD values. Laboratory duplicate RPD data were limited to LCS/LCSD and MS/MSD pairs and was acceptable. No field duplicates were identified. PCBs. All required holding times were met with one exception. Sample -049 was re-extracted 11 days after the holding time expired; however, given the persistent nature of the analyte, no data wee qualified as a result of the expired holding time. No method blank contamination was detected. LCS, MS, and surrogate-recoveries were within laboratory control limits. Laboratory duplicate RPDs were acceptable. No field duplicates were identified. Priority Pollutant Metals. All required holding times were met. No method blank contamination was detected with one exception. The zinc reporting limit for one prep batch was elevated from 1 mg/kg to 2.5 mg/kg due to possible laboratory contamination of the sample or the extract. No data were qualified on this basis, as the only sample associated with blank contained zinc levels greater than 20 times the elevated reporting limit. LCS recoveries were within laboratory control limits. The MS mercury recovery for prep batch 0100945 was less than the lower control limit, and the mercury duplicate for the same batch was outside the control limits. Mercury values for -071, -072, and -077 were therefore flagged "I" as estimated. The MS antimony recovery for prep batch 010846 was less than the lower control limit; however a laboratory note stated "Multiple analyses indicate the percent recovery is outside the control limits due to a matrix effect." Because Page F-9 **Flart Crowser** 15191-01 January 18, 2002

	the balance of the QC data for this batch was acceptable, the results were
	not qualified. The prep batch 0J29009 contained two matrix spikes. Arsenic
	and selenium recoveries for 0J2909-MS2 were below the lower control limits;
	however, the balance of the associated QC was acceptable, and no data
	were qualified. Laboratory duplicate data were otherwise acceptable. No
-	field duplicates were identified.
	TCID Land All garginal holding simon was now to the delant.
	TCLP Lead. All required holding times were met. No method blank contamination was reported. LCS and MS recoveries were acceptable. No
	•
	duplicate data were reported.
	Sample Prefix Nos. 5106-000925
	NCA ID No. P009762
	Six soil samples were submitted to the laboratory. The following analyses
	were performed on one or more samples:
	,
	■ TPH as Diesel/Oil (NWTPH-Dx); and
	■ PAHs (EPA Method 8270 SIM);
	This (LITT Method 0270 Shvi),
· · · · · · · · · · · · · · · · · · ·	TPH-as-Diesel/Oil. All required holding times-were met. No-method blank
	contamination was detected. Surrogate and LCS recoveries were within
	laboratory control limits. MS analysis is not required by the method, and
	none was performed. Laboratory duplicate RPDs were within the laboratory
-	control limits. No-field-duplicates were-identified.
	•
	PAHs. All required holding times were met. No method blank contamination
	was detected. LCS, MS, and surrogate recoveries were within laboratory
	control limits. Duplicate RPD data were acceptable. No field duplicates
	were identified.
Annondiy I	: Data Gap Investigation Soil Samples (October and November 2000)
Appelluix L.	Data Dap investigation don damples (October and November 2000)
	Sample Prefix Nos. 5106-001026, 5106-001027, and 5106-001030
	ESL ID No. 0010192
	Thirty soil samples were submitted to the laboratory, including some samples
	that were not analyzed in this set. The following analyses were performed on
	one or more samples:
	■ TPH Identification (NWTPH-HCID);
	■ TPH as Diesel/Oil (NWTPH-Dx);

		•
		
	- DTEV (FDA MARIA - 1 0020A).	
	■ BTEX, (EPA Method 8020A);	
<u> </u>	PAHs, (EPA-Method-8270-SIM); and	
	T. 14 (1) Urpa coog/zoog r. (-) A-(- 1)	
•	■ Total Arsenic and Lead (EPA 6000/7000 Series Methods).	
	TPH Identification. All required holding times were met. No method blank	
	contamination was detected. LCS and MS analyses are not required by	
	NWTPH-HCID procedure, and were not performed. Surrogate recoveries	
	were acceptable. Laboratory duplicate results were acceptable. No field	
<u> </u>	duplicates were identified.	
	duplicates were identified.	
· · · · · · · · · · · · · · · · · · ·	TPH as Diesel/Oil. All required holding times were met. No method blank	
	contamination was detected. No LCS analysis was performed. Surrogate	_
	recoveries were within laboratory control limits. MS recoveries were within	
	control limits. Laboratory duplicate RPDs were acceptable. No field	
	duplicates were identified.	
· N	PAHs. All required holding times were met. No method blank contamination	
	was detected. LCS and MS/MSD-recoveries-were-within-laboratory control	
•	limits. Laboratory duplicate RPDs were acceptable. Surrogate recoveries for	
	-067, -068, -085, and -086 were elevated. Since the bias was positive, and	
	all other associated QC was acceptable, no data were qualified based on	
	surrogate recoveries. No field duplicates were identified.	
	Total Arsenic and Lead. All required holding times were met. No method	
······································	blank contamination was detected. LCS and MS recoveries were within	***************************************
	laboratory control limits. Laboratory duplicate RPDs were acceptable with	
	the following exception. The RPD for the lead duplicate was 35.7 percent as	
	compared to a control limit of 20 percent. Since the RPD for the lead	· .
	MS/MSD pair was acceptable, no data were qualified. No field duplicates	
	were identified.	
	Sample Prefix Nos. 5106-001024 & 5106-001025	
	NCA ID No. P010845	
	Fifty-five soil samples were submitted to the laboratory, including some	
•	samples that were not analyzed in this set. The following analyses were	•
	performed on one or more samples:	
	- TRU as Diosal/Oil (NIMTRU Da).	
	■ TPH as Diesel/Oil (NWTPH-Dx);	•
	■ PAHs (EPA Method 8270 SIM); and	
-	TOCA (EDA Mothad 9260P)	
•	■ VOCs (EPA Method 8260B).	
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TPH as Diesel/Oil. All required holding times were met. No method blank contamination was detected. Surrogate and LCS recoveries were within laboratory control limits. MS analysis is not required by the method, and none was performed. Laboratory duplicate RPDs were within the laboratory control limits. The chain of custody specified that sample -047 be performed in duplicate; however, no duplicate analysis was reported in this sample set. No field duplicates were identified. PAHs. All required holding times were met. No method blank contamination was detected. LCS and surrogate recoveries were within laboratory control limits. MS/MSD results were not useable, as analyte concentrations in the native sample were greater than four times the spiking levels. Laboratory duplicate data were acceptable. No field duplicates were identified. VOCs. All required holding times were met. No method blank contamination was detected. LCS, MS, and surrogate recoveries were within laboratory control limits. Laboratory duplicate RPD data were limited to the MS/MSD pair, and were acceptable. No field duplicates were identified. Sample Prefix No. 5106-0011106 ESL ID No. 0011030 Ten soil samples were submitted to the laboratory, including one sample that was not analyzed in this set. The following analyses were performed on one or more samples: ■ TPH as Diesel/Oil (NWTPH-Dx); PAHs, (EPA Method 8270 SIM); and Total Arsenic and Lead (EPA 6000/7000 Series Methods). TPH as Diesel/Oil. All required holding times were met. No method blank contamination was detected. LCS recovery was 69.2 percent compared to the lower control limit of 70 percent. Because the MS recoveries were within control limits, no data were qualified. Surrogate recoveries were within laboratory control limits. Laboratory duplicate RPDs were acceptable. No field duplicates were identified. PAHs. All required holding times were met. No method blank contamination was detected. Surrogate, LCS and MS/MSD recoveries were within laboratory control limits. Laboratory duplicate RPDs were acceptable. No field duplicates were identified. Hart Crowser Page F-12 15191-01 January 18, 2002

	
-	Total Arsenic and Lead. All required holding times were met. No method
	blank contamination was detected. LCS and MS recoveries were within
	laboratory control limits. Laboratory duplicate RPDs were acceptable with
:	the following exception. The RPD for the lead duplicate was 24.6 percent as
	compared to a control limit of 20 percent. Since the RPD for the lead
	MS/MSD pair was acceptable, no data were qualified. No field duplicates
	were identified.
Appendix M	: Baseline Groundwater (February and March 2000)
	Sample Prefix No. 4876-000229
	OAL ID No.L15336
	Nine water samples were submitted to the laboratory. The following analyses
	were performed on one or more samples:
	■ PAHs (EPA Method 8270 SIM);
	■ Full List Semivolatile Organic Compounds (Semi-VOAs, EPA Method 8270);
	■ VOCs (EPA Method 8260); and
•	
	Priority Pollutant Dissolved Metals (Dissolved Metals, EPA
	200/6000/7000 Series Methods).
**	
 	PAHs. All required holding times were met. No method blank contamination
	was detected. Surrogate and LCS recoveries were within laboratory control
	limits. No MS data were reported. Laboratory duplicate RPDs were
	acceptable. No field duplicates were identified.
	Semi-VOAs. All required holding times were met. No method blank
	contamination was detected. Surrogate and LCS/LCSD recoveries were
	within control limits. Laboratory duplicate RPD data were limited to the
	LCS/LCSD pair and were acceptable. No field duplicates were identified.
	VOCs. All required holding-times were met. No method-blank
	contamination was detected. Surrogate, LCS, and MS/MSD recoveries were
•	within control limits. Laboratory duplicates RPD data were limited to the
	MS/MSD pair and were acceptable. No field duplicates were identified.
• .	
	Dissolved Metals. All required holding times were met. No method blank
	contamination was detected. LCS and MS recoveries were within laboratory
	control limits. Laboratory duplicate RPDs were acceptable. No field
. •	duplicates were identified.
·	

Appendix N:	B-38 Area Characterization Groundwater (March 2000)
	Sample Prefix No. 4876-000313
	OAL ID No.L15469
	Two water samples were submitted to the laboratory. The following analyses
	were performed on one or more samples:
· ·	■ PAHS (FPA Method 8270 SIM);
	■ Semi-VOAs (EPA 8270); and
	■ BTEX (EPA Method 8020A).
	PAHs. All required holding times were met. No method blank contamination
	was detected. Surrogate and LCS/LCSD recoveries were within laboratory
	control limits. No MS data were reported. Laboratory duplicate was limited to the LCS/LCSD pair, and the RPDs were acceptable. No field duplicates
	were analyzed.
·	W.O.C. alialyzed.
	Semi-VOAs. All required holding times were met. No method blank
	contamination was detected. Surrogate and I.CS/LCSD recoveries were
	within control limits. No MS data were reported. Laboratory duplicate RPD
	data were limited to the LCS/LCSD pair and were acceptable.
<u>·</u>	BTEX. All required holding times were met. No method blank contamination
	was detected. Surrogate recoveries were acceptable: LCS, MS, and duplicate
	data were reported before the sample set arrived at the laboratory. All BTEX
	data_were_therefore-rejected
	Sample Prefix No. 4876-000316
	OAL ID No.L15520
	Four water samples were submitted to the laboratory. The following analyses
	were performed on one or more samples:
	■ PAHS (EPA Method 8270 SIM); and
	■ BTEXs (EPA Method 8020A).
	a bleas (EFA Mediod 6020A).
	PAHs. All required holding times were met. No method blank
·	contamination was detected. Surrogate and LCS/LCD recoveries were
	within laboratory control limits. No MS data were reported. Laboratory
•	
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, 	
•	duplicate RPD data were limited to the LCS/LCSD pair and were
	acceptable. No field duplicates were identified.
•	
	BTEX. All required holding times were met. No method blank contamination
-,	was detected. Surrogate recoveries were acceptable. Laboratory duplicate
	data were acceptable. LCS and MS data were reported before the sample set
	arrived at the laboratory. All BTEX data were therefore rejected.
Appendix O	Supplemental Site Characterization Groundwater (September 2000)
	Sample Prefix Nos. 5106-000922 & 5106-000925
	NCA ID No. P009764
,	Seven water samples were submitted to the laboratory. The following
<u>·</u>	analyses-were performed on one or more samples:
	- DATE (FDA L) of Tooms (IV.)
	■ PAHs (EPA Method 8270 SIM);
	■ VOCs (EPA Method 8260B);
	 Priority Pollutant Metals (EPA 6000/7000 Series Methods); and
	■ Bis(2-Ethylhexyl)phthalate (DEHP, EPA-Method 8270).
	PAHs. All required holding times were met. No method blank contamination
	was detected. Surrogate and LCS/LCSD recoveries were within laboratory
	— control limits. Laboratory duplicate-RPD data were limited to the LCS/LCSD
	pair and were acceptable. No field duplicates were identified.
	VOCa All remained helding the consequent No. 11.1.1
	VOCs. All required holding times were met. No method blank
	contamination was detected. Surrogate, LCS/LCSD, and MS/MSD recoveries
	were within laboratory control limits. Laboratory duplicate RPD data were
	limited to LCS/LCSD and MS/MSD pairs and were acceptable. No field duplicates were identified.
	duplicates were identified.
	Priority Pollutant Metals. All required holding times were met. No method
	blank contamination was detected. LCS recoveries were within laboratory
	control limits. There were two matrix spikes associated with this prep batch.
	The first spiked sample was from this sample set, and all recoveries were
	within control limits except for copper, which had a recovery of 62 percent as
	compared to the lower control limit of 75 percent. However, the LCS
	recovery for copper was acceptable. The second matrix spike was not from
	this sample set, and generally had poor recoveries. No data were qualified
<u></u>	and sample self and Benefally had poor recoveries. 140 data were qualified
•	
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	· · · · · · · · · · · · · · · · · · ·
	based on MS recoveries. Laboratory duplicate results were acceptable. No
	field duplicates were identified.
	DEHP. All required holding times were met. No method blank
	contamination was reported. Surrogate and LCS/LCSD recoveries were
· .	acceptable. Only LCS/LCSD duplicate data were reported and were
	acceptable. No field duplicates were identified.
Appendix	P: Data Gap Investigation Groundwater (October and November 2000)
	Sample Prefix Nos. 5106-001026, 5106-001027, 5106-001030,
	and 5106-001024
	ESL ID No. 0010191
	Eight-water-samples were submitted to the laboratory. The following analyses
	were performed on one or more samples:
	- TRU Discolor NATRUED
	■ TPH as Diesel/Oil (NWTPH-Dx);
<u></u>	■ PAHs (EPA Method 8270 SIM);
	■ VOCs (EPA Method 8260B); and
	■ Bis(2-Ethylhexyl)phthalate (DEHP, EPA Method 8270).
	TPH as Diesel/Oil. All required holding times were met. No method blank
	contamination was detected. Surrogate, LCS/LCSD, and MS-recoveries-were
	within laboratory control limits. Laboratory duplicate RPD data were limited
	to the LCS/LCSD pair and were acceptable.
	PAHs. All required holding times were met. No method blank contamination
•	was detected. Surrogate and LCS/LCSD recoveries were within laboratory
	control limits. Laboratory duplicate RPD data were acceptable. No field
	duplicates were identified.
,	VOCs. All required holding times were met. No method blank
	contamination was detected. Sample –100 was identified as "TB" on the
	chain of custody, and could possibly be a trip blank; however, there was
	nothing in the data to confirm its status. Surrogate and MS/MSD
	recoveries were within laboratory control limits. Laboratory-duplicate RPD
	data were limited to the MS/MSD pair and were acceptable. No field
•	duplicates were identified.

	DEHP. All required holding times were met. No method blank
	contamination was reported. Surrogate and LCS/LCSD recoveries were
	acceptable. Laboratory duplicate RPD data were limited to the LCS/LCSD
	pair and were acceptable. No field duplicates were identified.
	Data Prefix No. 5106-001024
	NCA Lab ID No. P010847
	NCA Eab ID No. FU 10041
· · · · · · · · · · · · · · · · · · ·	Three water samples were submitted to the laboratory. The following
	analyses were performed on one or more samples:
	·
	■ TPH as Diesel/Oil (NWTPH-Dx);
	
•	■ PAHs (EPA Method 8270 SIM); and
	■ VOCs (EPA Method-8260B).
	TPH as Diesel/Oil. All required holding times were met. No method blank
	contamination was detected. Surrogate, LCS/LCSD, and MS recoveries were
·	within laboratory control limits. Laboratory duplicate-RPD data were limited to
	the LCS/LCSD pair and were acceptable. No field duplicates were identified.
····	PAHs. All required holding times were met. No method blank contamination
·-	was detected. Surrogate and LCS/LCSD recoveries were within laboratory
	control limits. Laboratory duplicate RPD data were limited to the LCS/LCSD
	data pair and were acceptable. No field duplicates were identified.
·	
	VOCs. All required holding times were met. No method blank
	contamination was detected. Sample = 100 was identified as "TB" on the
	chain-of-custody, and-could possibly be a trip-blank; however, there was
	nothing in the data to confirm its status. There are no data in the sample set
<u> </u>	corresponding to this sample. Surrogate and LCS/LCSD recoveries were
	within laboratory control limits. There were no MS data provided.
	Laboratory duplicate RPD data were limited to the LCS/LCSD pair and were
	acceptable. No field duplicates were identified.
	Data Brofix No. 5406 004025
	Data Prefix No. 5106-001025 NCA Lab ID No. P010848
	NCA Lab ID No. PU 10040
	One water samples was submitted to the laboratory. The following analyses
	were performed:
	■ TPH as Diesel/Oil (NWTPH-Dx);
<u> </u>	
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	■ PAHs (EPA Method 8270 SIM); and
	■ DEHP-(EPA-Method-8270C).
	= DEFIT (EL A Medioù 02/00).
	TPH as Diesel/Oil. All required holding times were met. No method blank
	contamination was detected. Surrogate and LCS/LCSD recoveries were
,	within laboratory control limits. MS data were acceptable. Laboratory
	duplicate data were acceptable. No field duplicates were identified.
	PAHs. All required holding times were met. No method blank contamination
	was detected. Surrogate and LCS/LCSD recoveries were within laboratory
	control-limits:No-MS-data-were-provided:Laboratory-duplicate-RPD-data
· · · · · · · · · · · · · · · · · · ·	were acceptable. No field duplicates were identified.
	DEHP. All required holding times were met. No method blank
<u> </u>	contamination was detected. Surrogate and LCS/LCSD recoveries were
•	within laboratory control limits. No MS data were provided. Laboratory
	duplicate RPD data were acceptable. No field duplicates were identified.
	The MS/MSD data reported for -020 was prepared/analyzed six days
	before the sample was prepped, and is therefore unacceptable. Data for
<u>. </u>	= 020 was rejected.
	Sample Prefix Nos. 5106-001106
	ESL ID No. 0011031
	Two water growler was a shorted to the lebentary The Cillerian and
	Two water samples were submitted to the laboratory. The following analyses
•	were performed on one or more samples:
	■ TPH as Diesel/Oil (NWTPH-Dx);
	■ PAHs (EPA Method 8270 SIM); and
	■ BNA SVOCs (EPA Method 8270C).
	TPH as Diesel/Oil. All required holding times-were met. No method blank
	contamination was detected. Surrogate, LCS, and MS recoveries were within
	laboratory control limits. Laboratory duplicate RPD data was limited to the
	LCS/LCSD pair, and was acceptable.
	DAMe All required holding times were man blameth at blad a control of
•	PAHs. All required holding times were met. No method blank contamination
	was detected. Surrogate and LCS/LCSD recoveries were within laboratory
	control limits. Laboratory duplicate RPD data was acceptable. No field duplicates were identified.
	ouplicates were lucitified.
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BNA SVOCs. All required holding times were met. No method blank contamination was detected. Surrogate and LCS/LCSD recoveries were within laboratory control limits. Laboratory duplicate RPD data was acceptable. No field duplicates were identified. MONITORING WELL INSTALLATION AND GROUNDWATER SAMPLING Appendix D: Laboratory Results and Chain-of-Custody Documentation: Soil Sample **NCA ID No. P1H0861** One soil sample was submitted to the laboratory. The following analyses were performed: TPH as Diesel/Oil (NWTPH-Dx); Total Metals (EPA 6000/7000 Series Methods); Volatile Organic Compounds (VOCs, EPA Method-8260A); and PAHs (EPA Method 8270M-SIM). TPH as Diesel/Oil. All required holding times were met. No method blank contamination was detected. Surrogate and LCS recoveries were within laboratory control limits with the following exception. The surrogate recovery for duplicate sample (1091079-DUP) was not recoverable due to sample dilution. No qualification was necessary. Laboratory duplicate RPDs were acceptable. Total Metals: All required holding times were met. No method blank contamination was detected. LCS and MS recoveries were within laboratory control limits with the following exceptions. Antimony MS recoveries of 51.1 and 46.9 percent-were less than control limits (75 to 125 percent); however, the laboratory notes state that these recoveries were outside of control limits due to sample dilution. Therefore, no qualification was necessary. The laboratory duplicate RPDs for antimony (44.4 percent), beryllium (43.8 percent), silver (40.3 percent), and thallium (62.9 percent) were greater than the control limit of 40 percent. However, since the original and duplicate sample results for these four metals were nondetect, the RPD is not applicable and no qualification was necessary. VOCs. All required holding times were met. No method blank contamination was detected. Surrogate, LCS, and MS recoveries were within laboratory control limits. MS/MSD RPDs were acceptable. Hart Crowser Page F-19 15191-01 January 18, 2002

	PAHs. All required holding times were met. No method blank contamination
	was detected. Surrogate, LCS, and MS recoveries were within laboratory
 	control limits with the following exception. The benzo(a)pyrene and pyrene
	MS spikes were not recoverable and the MS/MSD RPD was greater than
	— control limits due to a non-homogeneous sample matrix according to the
	laboratory. Additionally, the pyrene source result was more than four times
	greater than the pyrene spike level indicating that the percent recovery for
	pyrene is not applicable. Because the MS/MSD sample was from a separate
	sample batch, the remaining quality control samples were acceptable, and
	benzo(a)pyrene was not detected in the site sample, no qualification was
	necessary.
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* *	Laboratory Results and Chain-of-Custody Documentation: October 2001 Groundwater Monitoring Samples
	Clober 2001 Groundwater morntoring Samples
	NCA ID No. P1J0098
<u></u>	Four groundwater samples were submitted to the laboratory. The following
	analyses were performed on one or more of the samples:
	■ TPH as Diesel/Oil (NWTPH-Dx);
	■ Total and Dissolved Metals (EPA 6000/7000 Series Methods);
	■ VOCs (EPA Method 8260A);
	■ SVOCs (EPA Method 8270C); and
	PAHs (EPA Method 8270M-SIM).
	TPH as Diesel/Oil. All required holding times were met. No method blank
	contamination was detected. Surrogate and LCS/LCS Dup recoveries were
	within laboratory control limits. The LCS/LCS-Dup RPDs were within
	laboratory control limits.
	Total and Dissolved Metals. All required holding times were met. No
_ 	method blank contamination was detected. LCS and MS recoveries and
	duplicate RPDs were within laboratory control limits with the following
•	· ·
	exceptions. The dissolved metals duplicate RPDs for copper and lead
	exceptions. The dissolved metals duplicate RPDs for copper and lead exceeded the RPD limit of 20 percent. However, the original and duplicate
	exceeded the RPD limit of 20 percent. However, the original and duplicate
	exceeded the RPD limit of 20 percent. However, the original and duplicate sample results for both metals were nondetect indicating that the RPD criterion is not applicable and that no qualification is necessary.
	exceeded the RPD limit of 20 percent. However, the original and duplicate sample results for both metals were nondetect indicating that the RPD criterion is not applicable and that no qualification is necessary. VOCs. All required holding times were met. No method blank
	exceeded the RPD limit of 20 percent. However, the original and duplicate sample results for both metals were nondetect indicating that the RPD criterion is not applicable and that no qualification is necessary. VOCs. All required holding times were met. No method blank contamination was detected. Surrogate, LCS, and MS/MSD recoveries were
	exceeded the RPD limit of 20 percent. However, the original and duplicate sample results for both metals were nondetect indicating that the RPD criterion is not applicable and that no qualification is necessary. VOCs. All required holding times were met. No method blank

SVOCs. All required holding times were met. No method blank contamination was detected. Surrogate and LCS/LCS Dup recoveries were within laboratory control limits. LCS/LCS Dup RPDs were acceptable. PAHs. All-required holding times were met. No method blank contamination was detected. Surrogate and LCS/LCS Dup recoveries were within laboratory control limits. LCS/LCS Dup RPDs were acceptable. NCA ID No. P1J0097 Five groundwater samples were submitted to the laboratory. The following analyses'were performed on one or more of the samples: TPH as Diesel/Oil (NWTPH-Dx); Total and Dissolved Metals (EPA 6000/7000 Series Methods); VOCs (EPA Method 8260A); SVOCs (EPA Method 8270C); and PAHs (EPA Method 8270M-SIM). TPH as Diesel/Oil. All required holding times were met. No method blank contamination was detected. Surrogate and LCS/LCS Dup recoveries were within laboratory control limits. The LCS/LCS Dup RPDs were within laboratory control limits. The laboratory flagged the concentrations of diesel-range organics in samples 5106-011001-108 (MW-1) and 5106-01101-109 (MW-1 Dup) to indicate that the detected hydrocarbons have non-petroleum peaks suggesting the presence of biogenic interference. The results suggest the detected diesel-range organics may be an overestimate, if present in the sample. Total and Dissolved Metals. All required holding times were met. No method blank contamination was detected. LCS and MS recoveries and duplicate RPDs were within laboratory control limits. The dissolved metals duplicate RPDs for copper and lead exceeded the RPD limit of 20 percent. However, the original and duplicate sample results for both metals were nondetect indicating that the RPD criterion is not applicable and that no qualification is necessary. VOCs. All required holding times were met. No method blank contamination was detected. Surrogate, LCS, and MS/MSD recoveries were within laboratory control limits. MS/MSD RPDs were acceptable. Hart Crowser Page F-21

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SVOCs. All required holding times were met. No method blank contamination was detected. Surrogate and LCS/LCS Dup recoveries were within laboratory control limits. LCS/LCS Dup RPDs were acceptable. PAHs. All-required holding times were met. No method blank contamination was detected. Surrogate and LCS/LCS Dup recoveries were within laboratory control limits with the following exceptions. The laboratory flagged the surrogate recoveries for sample MW-1 Dup indicating that the surrogate recoveries exceeded control limits, the laboratory suspected a double spike of the surrogate solution during extraction, and that actual surrogate recoveries were believed to be one-half the reported values. The surrogate recoveries for sample MW-1 were within laboratory control limits supporting the laboratories claim that the surrogate solution was double spiked. Additionally, since no analytes were detected in sample MW-1 Dup, no qualification was necessary. Field Groundwater Duplicate Samples (MW-1/MW-1 Dup). The precision requirements for analyte detected in samples MW-1 and MW-1 Dup were met. All analytes detected were detected at concentrations less than five times their respective reporting limits. Therefore, a control limit of plus or minus the reporting limit was used to evaluate precision for these analytes. Equipment Blank. VOCs and bis(2-ethylhexyl)phthalate were not detected above method detection limits in sample 5106-011001-110 (Equipment Blank). Page F-22 Hart Crowser 15191-01 January 18, 2002